

The listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims

Claim 1 (canceled).

Claim 2 (currently amended): The method of Claim 14, wherein R¹⁵ is selected from H, ~~optionally substituted pyrrolidinyl, optionally substituted piperazinyl, optionally substituted piperidinyl, morpholinyl, 1,2,3,6-tetrahydro-pyridinyl, (optionally substituted pyrrolidinyl) C₁-C₂-alkyl, (optionally substituted piperidinyl) C₁-C₂-alkyl, (optionally substituted piperazinyl) C₁-C₂-alkyl, morpholinyl C₁-C₂-alkyl, C₁-C₄-alkylamino C₁-C₄-alkyl, C₁-C₄-hydroxyalkylamino, (optionally substituted pyrrolidinyl) C₁-C₂-alkylamino, (optionally substituted piperidinyl) C₁-C₂-alkylamino, (optionally substituted piperazinyl) C₁-C₂-alkylamino, morpholinyl C₁-C₂-alkylamino, optionally substituted pyrrolidinyl-C₁-C₄-alkoxy, optionally substituted azetidiny-C₁-C₄-alkoxy, tetrahydrofuryl-C₁-C₄-alkoxy, optionally substituted piperidinyl-C₁-C₄-alkoxy, C₁-C₄-alkylamino-C₁-C₄-alkoxy, tetrahydrofuryloxy, optionally substituted piperidinyl-oxy, and optionally substituted phenoxy, C₁-C₄-alkylaminocarbonyl and C₁-C₄-alkylaminethiocarbonyl; wherein R¹⁶ is selected from H, ~~5-6 membered nitrogen-containing heterocyclylcarbonyl, C₁-C₄-alkylaminocarbonyl, and C₁-C₄-alkylaminomethyl, and 5-6 membered nitrogen-containing heterocyclylmethyl~~; and wherein R¹⁷ is selected from halo, C₁-C₂-alkyl, ~~thienylsulfonyl- C₁-C₂-alkyl, optionally substituted 5-6 membered heteroarylsulfonyl C₁-C₂-alkyl, optionally substituted phenoxy, and C₃-C₆-cycloalkyl-C₂-C₄-alkynyl, and pharmaceutically acceptable derivatives thereof.~~~~

Claim 3 (currently amended): The method of Claim 2, wherein R¹⁵ is selected from H, tetrahydro-furanyloxy, 1-methylpyrrolidin-2-ylmethoxy, 2-pyrrolidinylmethoxy, 3-pyrrolidinylmethoxy, 1-Boc-pyrrolidin-2-ylmethoxy, 4-piperidinylmethoxy, 1-Boc-piperidin-4-ylmethoxy, 1-Boc-piperidin-4-ylethoxy, piperidin-4-ylethoxy, 1-methyl-piperidin-4-ylmethoxy, 1-Boc-azetidin-3-ylmethoxy, 1-methyl-azetidin-3-ylmethoxy, 3-azetidinylmethoxy, 1-methyl-piperidin-4-yloxy, ~~phenyloxy, phenoxy, 4-(pyrrolidin-1-ylmethyl)phenoxy, and dimethylaminoethoxy, 1-piperidinylmethyl, 1-(piperidin-1-yl)ethyl, 3-methylpiperidin-1-~~

~~ylmethyl, 1-pyrrolidinylmethyl, 2,2,6,6-tetramethylpiperidin-1-ylmethyl, 2,6-dimethylpiperidin-1-ylmethyl, dimethylaminomethyl, diethylaminomethyl, diethylaminothiocarbonyl, diethylaminocarbonyl, N-Boc-N-isopropylaminomethyl, isopropylaminomethyl, 2-thienylsulfonylmethyl, hydroxypropylamino, 4-ethyl-piperidin-1-yl, 4-(2-pyridyl)piperidin-1-yl, 1-methylpiperidin-4-yl, 4-(2-pyrazinyl)piperidin-1-yl, 1-methyl-1,2,3,6-tetrahydro-pyridin-4-yl, 1,2,3,6-tetrahydro-pyridin-4-yl, and 1-Boc-1,2,3,6-tetrahydro-pyridin-4-yl; wherein R¹⁶ is selected from H, 1-piperidinylcarbonyl, diethylaminocarbonyl, and diethylaminomethyl, 1-piperidinylmethyl; and wherein R¹⁷ is selected from chloro, bromo, methyl and cyclopropylethynyl, and pharmaceutically acceptable derivatives thereof.~~

Claim 4 (currently amended): The method of Claim 3, wherein R¹⁷ is chloro or bromo, ~~and pharmaceutically acceptable derivatives thereof.~~

Claim 5 (currently amended): The method of Claim 14, wherein R¹⁵ is selected from H, ~~optionally substituted pyrrolidinyl, optionally substituted piperazinyl, optionally substituted piperidinyl, morpholinyl, 1,2,3,6-tetrahydro-pyridinyl, (optionally substituted pyrrolidinyl)-C₁-C₂-alkyl, (optionally substituted piperidinyl)-C₁-C₂-alkyl, (optionally substituted piperazinyl)-C₁-C₂-alkyl, morpholinyl-C₁-C₂-alkyl, (optionally substituted pyrrolidinyl)-C₁-C₂-alkylamino, (optionally substituted piperidinyl)-C₁-C₂-alkylamino, (optionally substituted piperazinyl)-C₁-C₂-alkylamino, morpholinyl-C₁-C₂-alkylamino, C₁-C₄-alkylamino-C₁-C₄-alkyl, C₁-C₄-hydroxyalkylamino, optionally substituted pyrrolidinyl-C₁-C₄-alkoxy, optionally substituted azetidiny-C₁-C₄-alkoxy, tetrahydrofuryl-C₁-C₄-alkoxy, optionally substituted piperidinyl-C₁-C₄-alkoxy, C₁-C₄-alkylamino-C₁-C₄-alkoxy, tetrahydrofuryloxy, optionally substituted piperidinyl, and optionally substituted phenoxy, C₁-C₄-alkylaminocarbonyl and C₁-C₄-alkylaminocarbonyl; wherein R¹⁶ is selected from H, 5-6-membered nitrogen-containing heterocyclylcarbonyl, C₁-C₄-alkylaminocarbonyl, and C₁-C₄-alkylaminomethyl, and 5-6-membered nitrogen-containing heterocyclylmethyl; and wherein R¹⁷ is selected from C₃-C₆-cycloalkyl and phenyl optionally substituted with one or two substituents selected from halo, C₁-C₄-alkylamino, amino, nitro, C₁-C₄-alkoxy, C₁-C₂-haloalkyl, hydroxy, C₁-C₄-alkylthio, C₁-C₄-alkylcarbonylamino, (optionally substituted phenyl)sulfonylamino, cyano, C₁-C₂-haloalkoxy, 5- or 6-membered N-containing heterocyclyl, aminocarbonyl,~~

~~(6-membered N-containing heterocyclyl)sulfonyl, C₁-C₂-haloalkylcarbamoylamino~~
~~sulfonyl, and (optionally substituted phenyl)amino~~
~~sulfonyl, and pharmaceutically acceptable derivatives thereof.~~

Claim 6 (currently amended): The method of Claim 5, wherein R¹⁵ is selected from H, tetrahydro-furanyloxy, 1-methylpyrrolidin-2-ylmethoxy, 2-pyrrolidinylmethoxy, 3-pyrrolidinylmethoxy, 1-Boc-pyrrolidin-2-ylmethoxy, 4-piperidinylmethoxy, 1-Boc-piperidin-4-ylmethoxy, 1-Boc-piperidin-4-ylethoxy, piperidin-4-ylethoxy, 1-methyl-piperidin-4-ylmethoxy, 1-Boc-azetidin-3-ylmethoxy, 1-methyl-azetidin-3-ylmethoxy, 3-azetidinylmethoxy, 1-methyl-piperidin-4-yloxy, ~~phenyloxy, phenoxy~~ 4-(pyrrolidin-1-ylmethyl)phenoxy, and dimethylaminoethoxy, ~~1-piperidinylmethyl, 1-(piperidin-1-yl)ethyl, 3-methylpiperidin-1-ylmethyl, 1-pyrrolidinylmethyl, 2,2,6,6-tetramethylpiperidin-1-ylmethyl, 2,6-dimethylpiperidin-1-ylmethyl, dimethylaminomethyl, diethylaminomethyl, diethylaminothiocarbonyl, diethylaminocarbonyl, N-Boc-N-isopropylaminomethyl, isopropylaminomethyl, 2-thienylsulfonylmethyl, hydroxypropylamino, 4-ethyl-piperidin-1-yl, 4-(2-pyridyl)piperidin-1-yl, 1-methylpiperidin-4-yl, 4-(2-pyrazinyl)piperidin-1-yl, 1-methyl-1,2,3,6-tetrahydro-pyridin-4-yl, 1,2,3,6-tetrahydro-pyridin-4-yl, and 1-Boc-1,2,3,6-tetrahydro-pyridin-4-yl;~~ wherein R¹⁶ is selected from H, ~~1-piperidinylearbonyl,~~ diethylaminocarbonyl, and diethylaminomethyl, ~~1-piperidinylmethyl;~~ and wherein R¹⁷ is selected from cyclopropyl and phenyl-~~optionally substituted with aminosulfonyl, and pharmaceutically acceptable derivatives thereof.~~

Claim 7 (currently amended): The method of Claim 6, wherein R¹⁷ is unsubstituted phenyl; ~~and pharmaceutically acceptable derivatives thereof.~~

Claim 8 (currently amended): The method of Claim 14, wherein R¹⁵ is selected from H, ~~optionally substituted pyrrolidinyl, optionally substituted piperazinyl, optionally substituted piperidinyl, morpholinyl, 1,2,3,6-tetrahydro-pyridinyl, (optionally substituted pyrrolidinyl) C₁-C₂-alkyl, (optionally substituted piperidinyl) C₁-C₂-alkyl, (optionally substituted piperazinyl) C₁-C₂-alkyl, morpholinyl C₁-C₂-alkyl, (optionally substituted pyrrolidinyl) C₁-C₂-alkylamino, (optionally substituted piperidinyl) C₁-C₂-alkylamino, (optionally substituted piperazinyl) C₁-C₂-alkylamino, morpholinyl C₁-C₂-alkylamino, C₁-C₄-alkylamino C₁-C₄-alkyl, C₁-C₄-hydroxyalkylamino,~~ optionally substituted pyrrolidinyl-C₁-C₄-alkoxy, optionally substituted

azetidiny-C₁-C₄-alkoxy, tetrahydrofuryl-C₁-C₄-alkoxy, optionally substituted piperidiny-C₁-C₄-alkoxy, C₁-C₄-alkylamino-C₁-C₄-alkoxy, tetrahydrofuryloxy, optionally substituted piperidinyloxy, and optionally substituted phenoxy, ~~C₁-C₄-alkylaminocarbonyl and C₁-C₄-alkylaminothiocarbonyl~~; wherein R¹⁶ is selected from H, ~~5-6 membered nitrogen-containing heterocyclylcarbonyl~~, C₁-C₄-alkylaminocarbonyl, and C₁-C₄-alkylaminomethyl, ~~and 5-6 membered nitrogen-containing heterocyclylmethyl~~; and wherein R¹⁷ is selected from ~~optionally substituted indazolyl, optionally substituted indolyl, unsubstituted 5-membered oxygen or sulfur containing heterocaryl~~, unsubstituted thienyl, unsubstituted 6-membered nitrogen-containing heterocyclyl, and 6-membered nitrogen-containing heterocyclyl substituted with ~~one or more substituents independently selected from pyridyl, phenyl,~~

~~C₁-C₄-alkyl, C₁-C₂-haloalkyl, C₁-C₂ alkoxy, amino, halo, piperidinyl, morpholinyl, C₁-C₂-alkylpiperazinyl, C₁-C₂-alkylaminothiocarbonyl, N,N-di-C₁-C₂-alkylamino-C₁-C₄-alkylenyl, N-C₁-C₂-alkylamino-C₁-C₄-alkylenyl, morpholinyl-C₁-C₄-alkylenylaminocarbonyl, aminocarbonyl, C₁-C₂-haloalkylcarbonylamino, morpholinyl-C₁-C₄-alkylenylamino, N,N-di-C₁-C₂-alkylamino and N,N-di-C₁-C₂-alkylamino-C₁-C₄-alkylenylamino;~~
~~and pharmaceutically acceptable derivatives thereof.~~

Claim 9 (currently amended): The method of Claim 8, wherein R¹⁵ is selected from ~~H,~~ tetrahydro-furanyloxy, 1-methylpyrrolidin-2-ylmethoxy, 2-pyrrolidinylmethoxy, 3-pyrrolidinylmethoxy, 1-Boc-pyrrolidin-2-ylmethoxy, 4-piperidinylmethoxy, 1-Boc-piperidin-4-ylmethoxy, 1-Boc-piperidin-4-ylethoxy, piperidin-4-ylethoxy, 1-methyl-piperidin-4-ylmethoxy, 1-Boc-azetidin-3-ylmethoxy, 1-methyl-azetidin-3-ylmethoxy, 3-azetidinyloxy, 1-methyl-piperidin-4-yloxy, phenoxy, 4-(pyrrolidin-1-ylmethyl)phenoxy, and dimethylaminoethoxy, ~~1-piperidinylmethyl, 1-(piperidin-1-yl)ethyl, 3-methylpiperidin-1-ylmethyl, 1-pyrrolidinylmethyl, 2,2,6,6-tetramethylpiperidin-1-ylmethyl, 2,6-dimethylpiperidin-1-ylmethyl, dimethylaminomethyl, diethylaminomethyl, diethylaminothiocarbonyl, diethylaminocarbonyl, N-Boc-N-isopropylaminomethyl, isopropylaminomethyl, 2-thienylsulfonylmethyl, hydroxypropylamino, 4-ethylpiperidin-1-yl, 4-(2-pyridyl)piperidin-1-yl, 1-methylpiperidin-4-yl, 4-(2-pyrazinyl)piperidin-1-yl, 1-methyl-1,2,3,6-tetrahydro-pyridin-4-yl, 1,2,3,6-tetrahydro-pyridin-4-yl, and 1-Boc-1,2,3,6-tetrahydro-pyridin-4-yl~~; wherein R¹⁶ is selected from H, ~~4-piperidinylcarbonyl, diethylaminocarbonyl, and diethylaminomethyl, 1-piperidinylmethyl; and~~

wherein R¹⁷ is selected from ~~5-indazolyl, 1-Boc-indol-5-yl, unsubstituted thienyl, 5-tert-butylloxazol-2-yl and 4-pyridyl substituted with one or more substituents independently selected from methoxy and chloro, and pharmaceutically acceptable derivatives thereof.~~

Claim 10 (currently amended): The method of Claim ~~9~~ 8, wherein R¹⁷ is 4-pyridyl; ~~and pharmaceutically acceptable derivatives thereof.~~

Claim 11 (currently amended): The method of Claim 14 ~~and pharmaceutically acceptable derivatives thereof, wherein the compound is selected from:~~

~~1-[6-(3-Methyl-piperidin-1-ylmethyl)-pyridin-2-yl]-3-(2-phenyl-thiazol-4-yl)-urea;~~
~~1-[4-(Piperidine-1-carbonyl)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;~~
~~1-(2-Chloro-thiazol-4-yl)-3-[4-(piperidine-1-carbonyl)-pyridin-2-yl]-urea;~~
~~N,N-Diethyl-2-[3-(2-pyridin-4-yl-thiazol-4-yl)-ureido]-isonicotinamide;~~
~~N,N-Diethyl-2-[3-(2-phenyl-thiazol-4-yl)-ureido]-isonicotinamide;~~
~~2-[3-(2-Bromo-thiazol-4-yl)-ureido]-N,N-diethyl-isonicotinamide;~~
~~1-(4-Diethylaminomethyl-pyridin-2-yl)-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;~~
~~1-[6-(2,6-Dimethyl-piperidin-1-ylmethyl)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;~~
~~1-[6-(1-Piperidin-1-yl-ethyl)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;~~
~~2-[(6-[3-(2-Pyridin-4-yl-thiazol-4-yl)-ureido]-pyridin-2-ylamino)-methyl]-piperidine-1-carboxylic acid tert-butyl ester;~~
~~1-[6-[(Piperidin-2-ylmethyl)-amino]-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;~~
~~(S)-1-[6-(3-Methyl-piperidin-1-ylmethyl)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;~~
~~(R)-1-[6-(3-Methyl-piperidin-1-ylmethyl)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;~~
~~1-(2-Chloro-thiazol-4-yl)-3-(6-piperidin-1-ylmethyl-pyridin-2-yl)-urea;~~
1-(2-Bromo-thiazol-4-yl)-3-[6-(2-piperidin-4-yl-ethoxy)-pyridin-2-yl]-urea;
1-(2-Chloro-thiazol-4-yl)-3-[6-(2-piperidin-4-yl-ethoxy)-pyridin-2-yl]-urea;
1-[6-(Azetidin-3-ylmethoxy)-pyridin-2-yl]-3-(2-bromo-thiazol-4-yl)-urea;
1-[6-(Azetidin-3-ylmethoxy)-pyridin-2-yl]-3-(2-chloro-thiazol-4-yl)-urea;
1-(2-Bromo-thiazol-4-yl)-3-[6-(piperidin-4-ylmethoxy)-pyridin-2-yl]-urea;
1-(2-Chloro-thiazol-4-yl)-3-[6-(piperidin-4-ylmethoxy)-pyridin-2-yl]-urea;

tert-Butyl 3-{6-[3-(2-pyridin-4-yl-thiazol-4-yl)-ureido]-pyridin-2-yloxymethyl}-pyrrolidine-1-carboxylate;

1-(2-Pyridin-4-yl-thiazol-4-yl)-3-[6-(pyrrolidin-3-ylmethoxy)-pyridin-2-yl]-urea;

1-(2-Cyclopropyl-thiazol-4-yl)-3-[6-(2-piperidin-4-yl-ethoxy)-pyridin-2-yl]-urea;

~~1-[6-(Isopropylamino-methyl)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;~~

~~1-[6-(Isopropylamino-methyl)-pyridin-2-yl]-3-(2-phenyl-thiazol-4-yl)-urea;~~

~~1-(2-Bromo-thiazol-4-yl)-3-[6-(isopropylamino-methyl)-pyridin-2-yl]-urea;~~

1-(2-Bromo-thiazol-4-yl)-3-[6-(1-methyl-pyrrolidin-2-ylmethoxy)-pyridin-2-yl]-urea;

1-(2-Chloro-thiazol-4-yl)-3-[6-(1-methyl-pyrrolidin-2-ylmethoxy)-pyridin-2-yl]-urea;

1-(2-phenylthiazol-4-yl)-3-(6-*p*-pyrrolidin-1-ylmethylphenoxy)pyridin-2-yl)urea;

1-(2-Pyridin-4-yl-thiazol-4-yl)-3-[6-(tetrahydro-furan-3-yloxy)-pyridin-2-yl]-urea;

~~1-[2-(1H-Indazol-5-yl)-thiazol-4-yl]-3-(6-piperidin-1-ylmethyl-pyridin-2-yl)-urea;~~

~~1-(1'-Methyl-1',2',3',6'-tetrahydro-[2,4']bipyridinyl-6-yl)-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;~~

~~1-(2-Bromo-thiazol-4-yl)-3-(1'-methyl-1',2',3',6'-tetrahydro-[2,4']bipyridinyl-6-yl)-urea;~~

~~1-(1'-Methyl-1',2',3',6'-tetrahydro-2[2,4']bipyridinyl-6-yl)-3-(2-phenyl-thiazol-4-yl)-urea;~~

~~1-[6-(3-Hydroxy-propylamino)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;~~

~~1-(2-Bromo-thiazol-4-yl)-3-[6-(3-hydroxy-propylamino)-pyridin-2-yl]-urea;~~

~~1-(1'-Methyl-1',2',3',4',5',6'-hexahydro-[2,4']bipyridinyl-6-yl)-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;~~

~~1-(1'-Methyl-1',2',3',4',5',6'-hexahydro-[2,4']bipyridinyl-6-yl)-3-(2-phenyl-thiazol-4-yl)-urea;~~

~~6-[3-(2-Pyridin-4-yl-thiazol-4-yl)-ureido]-3',6'-dihydro-2'H-[2,4']bipyridinyl-1'-carboxylic acid
tert-butylester;~~

~~1-(2-Pyridin-4-yl-thiazol-4-yl)-3-(1',2',3',6'-tetrahydro-[2,4']bipyridinyl-6-yl)-urea;~~

1-(2-Pyridin-4-yl-thiazol-4-yl)-3-[6-(tetrahydro-furan-3-ylmethoxy)-pyridin-2-yl]-urea;

2-[6-[3-(2-Pyridin-4-yl-thiazol-4-yl)-ureido]-pyridin-2-yloxymethyl]-pyrrolidine-1-carboxylic
acid tert-butyl ester;

1-(2-Pyridin-4-yl-thiazol-4-yl)-3-[6-(pyrrolidin-2-ylmethoxy)-pyridin-2-yl]-urea;

~~6-[3-(2-Pyridin-4-yl-thiazol-4-yl)-ureido]-pyridine-2-carbothioic acid diethylamide;~~

~~1-(2-Bromo-thiazol-4-yl)-3-[6-(3-methyl-piperidin-1-ylmethyl)-pyridin-2-yl]-urea;~~

~~1-(2-Chloro-thiazol-4-yl)-3-[6-(3-methyl-piperidin-1-ylmethyl)-pyridin-2-yl]-urea;~~

~~1-(2-Phenyl-thiazol-4-yl)-3-[4-(piperidine-1-carbonyl)-pyridin-2-yl]-urea;~~

~~1-(2-Bromo-thiazol-4-yl)-3-[4-(piperidine-1-carbonyl)-pyridin-2-yl]-urea;~~

1-[2-(2-Methoxy-pyridin-4-yl)-thiazol-4-yl]-3-(6-phenoxy-pyridin-2-yl)-urea;
1-[2-(2-Methoxy-pyridin-4-yl)-thiazol-4-yl]-3-[6-(1-methyl-pyrrolidin-2-ylmethoxy)-pyridin-2-yl]-urea;
1-[6-(2-Dimethylamino-ethoxy)-pyridin-2-yl]-3-[2-(2-methoxy-pyridin-4-yl)-thiazol-4-yl]-urea;
1-[6-(1-Methyl-pyrrolidin-2-ylmethoxy)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;
~~1-(2-phenylthiazol-4-yl)-3-(6-pyrrolidin-1-ylmethyl-pyridin-2-yl)urea;~~
~~1-(6-Diethylaminomethylpyridin-2-yl)-3-(2-phenylthiazol-4-yl)urea;~~
(S)-1-[6-(1-Methylpyrrolidin-2-ylmethoxy)pyridin-2-yl]-3-(2-phenylthiazol-4-yl)urea;
1-[6-(2-Piperidin-4-yl-ethoxy)pyridin-2-yl]-3-[2-phenylthiazol-4-yl]urea;
~~1-[6-(4-Ethylpiperazin-1-yl)-pyridin-2-yl]-3-(2-phenylthiazol-4-yl)urea;~~
~~Diethyl 6-[3-(2-phenylthiazol-4-yl)ureido]pyridine-2-carboxamide;~~
1-(2-Pyridin-4-yl-thiazol-4-yl)-3-(6-*p*-pyrrolidin-1-ylmethylphenoxy)pyridin-2-yl)urea;
1-(2-Bromothiazol-4-yl)-3-(6-*p*-pyrrolidin-1-ylmethylphenoxy)pyridin-2-yl)urea;
1-[6-(Piperidin-4-ylmethoxy)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;
1-[6-(1-Methyl-piperidin-4-ylmethoxy)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;
1-[6-(1-Methyl-piperidin-4-yloxy)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;
1-[6-(1-Methyl-azetidin-3-ylmethoxy)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;
1-[6-(Azetidin-3-ylmethoxy)-pyridin-2-yl]-3-(2-phenyl-thiazol-4-yl)-urea;
1-[6-(1-Methyl-azetidin-3-ylmethoxy)-pyridin-2-yl]-3-(2-phenyl-thiazol-4-yl)-urea;
1-(2-Phenyl-thiazol-4-yl)-3-[6-(piperidin-4-ylmethoxy)-pyridin-2-yl]-urea;
1-[6-(1-Methyl-piperidin-4-ylmethoxy)-pyridin-2-yl]-3-(2-phenyl-thiazol-4-yl)-urea;
1-[6-(1-Methyl-piperidin-4-yloxy)-pyridin-2-yl]-3-(2-phenyl-thiazol-4-yl)-urea;
1-[6-(2-Piperidin-4-yl-ethoxy)-pyridin-2-yl]-3-(2-thiophen-2-yl-thiazol-4-yl)-urea; and
1-[6-(1-Methyl-pyrrolidin-2-ylmethoxy)-pyridin-2-yl]-3-[2-(thiophene-2-sulfonylmethyl)-thiazol-4-yl]-urea;
~~1-[2-(2-Methoxy-pyridin-4-yl)-thiazol-4-yl]-3-(6-piperidin-1-ylmethyl-pyridin-2-yl)-urea; and~~
~~[2-(2-Chloro-pyridin-4-yl)-thiazol-4-yl]-3-(6-piperidin-1-ylmethyl-pyridin-2-yl)-urea;~~
and pharmaceutically acceptable salts thereof.

Claim 12 (currently amended): The method of Claim 14 ~~and pharmaceutically acceptable derivatives thereof~~, wherein the compound is selected from:

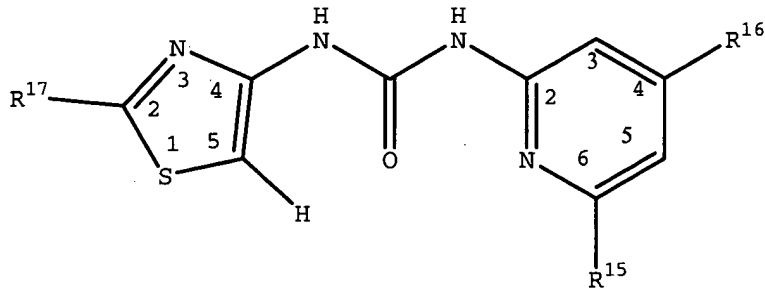
~~1-[6-(3-Methyl-piperidin-1-ylmethyl)-pyridin-2-yl]-3-(2-phenyl-thiazol-4-yl)-urea;~~
~~1-[4-(Piperidine-1-carbonyl)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;~~
~~N,N-Diethyl-2-[3-(2-pyridin-4-yl-thiazol-4-yl)-ureido]-isonicotinamide;~~
~~1-(4-Diethylaminomethyl-pyridin-2-yl)-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;~~
~~1-[6-(2,6-Dimethyl-piperidin-1-ylmethyl)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;~~
~~1-[6-(1-Piperidin-1-yl-ethyl)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;~~
~~2-((6-[3-(2-Pyridin-4-yl-thiazol-4-yl)-ureido]-pyridin-2-ylamino)-methyl)-piperidine-1-carboxylic acid tert-butyl ester;~~
~~1-[6-((Piperidin-2-ylmethyl)-amino)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;~~
~~(S)-1-[6-(3-Methyl-piperidin-1-ylmethyl)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;~~
~~(R)-1-[6-(3-Methyl-piperidin-1-ylmethyl)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;~~
~~1-(2-Chloro-thiazol-4-yl)-3-(6-piperidin-1-ylmethyl-pyridin-2-yl)-urea;~~
1-(2-Bromo-thiazol-4-yl)-3-[6-(2-piperidin-4-yl-ethoxy)-pyridin-2-yl]-urea;
1-(2-Chloro-thiazol-4-yl)-3-[6-(2-piperidin-4-yl-ethoxy)-pyridin-2-yl]-urea;
1-[6-(Azetidin-3-ylmethoxy)-pyridin-2-yl]-3-(2-bromo-thiazol-4-yl)-urea;
1-[6-(Azetidin-3-ylmethoxy)-pyridin-2-yl]-3-(2-chloro-thiazol-4-yl)-urea;
1-(2-Bromo-thiazol-4-yl)-3-[6-(piperidin-4-ylmethoxy)-pyridin-2-yl]-urea;
1-(2-Chloro-thiazol-4-yl)-3-[6-(piperidin-4-ylmethoxy)-pyridin-2-yl]-urea;
3-(4-{3-[6-(1-Methyl-pyrrolidin-2-ylmethoxy)-pyridin-2-yl]-ureido}-thiazol-2-yl)-benzenesulfonamide;
tert-Butyl 3-{6-[3-(2-pyridin-4-yl-thiazol-4-yl)-ureido]-pyridin-2-yloxymethyl}-pyrrolidine-1-carboxylate;
1-(2-Pyridin-4-yl-thiazol-4-yl)-3-[6-(pyrrolidin-3-ylmethoxy)-pyridin-2-yl]-urea;
1-(2-Cyclopropyl-thiazol-4-yl)-3-[6-(2-piperidin-4-yl-ethoxy)-pyridin-2-yl]-urea;
~~Isopropyl-{6-[3-(2-pyridin-4-yl-thiazol-4-yl)-ureido]-pyridin-2-ylmethyl}-carbamic acid tert-butyl ester;~~
~~1-[6-(Isopropylamino-methyl)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;~~
~~Isopropyl-{6-[3-(2-phenyl-thiazol-4-yl)-ureido]-pyridin-2-ylmethyl}-carbamic acid tert-butyl ester;~~
~~1-[6-(Isopropylamino-methyl)-pyridin-2-yl]-3-(2-phenyl-thiazol-4-yl)-urea;~~
1-(2-Bromo-thiazol-4-yl)-3-[6-(1-methyl-pyrrolidin-2-ylmethoxy)-pyridin-2-yl]-urea;
1-(2-Chloro-thiazol-4-yl)-3-[6-(1-methyl-pyrrolidin-2-ylmethoxy)-pyridin-2-yl]-urea;

1-(2-phenylthiazol-4-yl)-3-(6-*p*-pyrrolidin-1-ylmethylphenoxy)pyridin-2-yl)urea;
1-(2-Pyridin-4-yl-thiazol-4-yl)-3-[6-(tetrahydro-furan-3-yloxy)-pyridin-2-yl]-urea;
~~1-[2-(1H-Indazol-5-yl)-thiazol-4-yl]-3-(6-piperidin-1-ylmethyl-pyridin-2-yl)-urea;~~
~~1-(1'-Methyl-1',2',3',6'-tetrahydro-[2,4']bipyridinyl-6-yl)-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;~~
~~1-(2-Bromo-thiazol-4-yl)-3-(1'-methyl-1',2',3',6'-tetrahydro-[2,4']bipyridinyl-6-yl)-urea;~~
~~1-(1'-Methyl-1',2',3',6'-tetrahydro-2[2,4']bipyridinyl-6-yl)-3-(2-phenyl-thiazol-4-yl)-urea;~~
~~1-[6-(3-Hydroxy-propylamino)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;~~
~~1-(2-Bromo-thiazol-4-yl)-3-[6-(3-hydroxy-propylamino)-pyridin-2-yl]-urea;~~
~~1-(1'-Methyl-1',2',3',4',5',6'-hexahydro-[2,4']bipyridinyl-6-yl)-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;~~
~~1-(1'-Methyl-1',2',3',4',5',6'-hexahydro-[2,4']bipyridinyl-6-yl)-3-(2-phenyl-thiazol-4-yl)-urea;~~
~~6-[3-(2-Pyridin-4-yl-thiazol-4-yl)-ureido]-3',6'-dihydro-2'H-[2,4']bipyridinyl-1'-carboxylic acid tert-butylester;~~
~~1-(2-Pyridin-4-yl-thiazol-4-yl)-3-(1',2',3',6'-tetrahydro-[2,4']bipyridinyl-6-yl)-urea;~~
1-(2-Pyridin-4-yl-thiazol-4-yl)-3-[6-(tetrahydro-furan-3-ylmethoxy)-pyridin-2-yl]-urea;
2-[6-[3-(2-Pyridin-4-yl-thiazol-4-yl)-ureido]-pyridin-2-yloxymethyl]-pyrrolidine-1-carboxylic acid tert-butyl ester;
1-(2-Pyridin-4-yl-thiazol-4-yl)-3-[6-(pyrrolidin-2-ylmethoxy)-pyridin-2-yl]-urea;
~~6-[3-(2-Pyridin-4-yl-thiazol-4-yl)-ureido]-pyridine-2-carboxylic acid diethylamide;~~
~~1-(2-Bromo-thiazol-4-yl)-3-[6-(3-methyl-piperidin-1-ylmethyl)-pyridin-2-yl]-urea;~~
~~1-(2-Chloro-thiazol-4-yl)-3-[6-(3-methyl-piperidin-1-ylmethyl)-pyridin-2-yl]-urea;~~
1-[2-(2-Methoxy-pyridin-4-yl)-thiazol-4-yl]-3-[6-(1-methyl-pyrrolidin-2-ylmethoxy)-pyridin-2-yl]-urea;
1-[6-(2-Dimethylamino-ethoxy)-pyridin-2-yl]-3-[2-(2-methoxy-pyridin-4-yl)-thiazol-4-yl]-urea;
1-[6-(1-Methyl-pyrrolidin-2-ylmethoxy)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;
~~1-(2-phenylthiazol-4-yl)-3-(6-pyrrolidin-1-ylmethyl-pyridin-2-yl)urea;~~
~~1-(6-Diethylaminomethylpyridin-2-yl)-3-(2-phenylthiazol-4-yl)urea;~~
(S)-1-[6-(1-Methylpyrrolidin-2-ylmethoxy)pyridin-2-yl]-3-(2-phenylthiazol-4-yl)urea;
1-[6-(2-Piperidin-4-yl-ethoxy)pyridin-2-yl]-3-[2-phenylthiazol-4-yl]urea;
~~1-[6-(4-Ethylpiperazin-1-yl)-pyridin-2-yl]-3-(2-phenylthiazol-4-yl)urea;~~
~~1-(2-phenylthiazol-4-yl)-3-[6-(4-pyrimidin-2-yl-piperazin-1-yl)pyridin-2-yl]urea;~~
~~Diethyl 6-[3-(2-phenylthiazol-4-yl)-ureido]-pyridine-2-carboxamide;~~

1-(2-Pyridin-4-yl-thiazol-4-yl)-3-(6-*p*-pyrrolidin-1-ylmethylphenoxy)pyridin-2-yl)urea;
 1-(2-Bromothiazol-4-yl)-3-(6-*p*-pyrrolidin-1-ylmethylphenoxy)pyridin-2-yl)urea;
 1-[6-(Piperidin-4-ylmethoxy)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;
 1-[6-(1-Methyl-piperidin-4-ylmethoxy)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;
 1-[6-(1-Methyl-piperidin-4-yloxy)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;
 1-[6-(1-Methyl-azetidin-3-ylmethoxy)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;
 1-[6-(Azetidin-3-ylmethoxy)-pyridin-2-yl]-3-(2-phenyl-thiazol-4-yl)-urea;
 1-[6-(1-Methyl-azetidin-3-ylmethoxy)-pyridin-2-yl]-3-(2-phenyl-thiazol-4-yl)-urea;
 1-(2-Phenyl-thiazol-4-yl)-3-[6-(piperidin-4-ylmethoxy)-pyridin-2-yl]-urea;
 1-[6-(1-Methyl-piperidin-4-ylmethoxy)-pyridin-2-yl]-3-(2-phenyl-thiazol-4-yl)-urea;
 1-[6-(1-Methyl-piperidin-4-yloxy)-pyridin-2-yl]-3-(2-phenyl-thiazol-4-yl)-urea;
 1-[6-(2-Piperidin-4-yl-ethoxy)-pyridin-2-yl]-3-(2-thiophen-2-yl-thiazol-4-yl)-urea; and
 1-[6-(1-Methyl-pyrrolidin-2-ylmethoxy)-pyridin-2-yl]-3-[2-(thiophene-2-sulfonylmethyl)-thiazol-4-yl]-urea;
~~1-[2-(2-Methoxy-pyridin-4-yl)-thiazol-4-yl]-3-(6-piperidin-1-ylmethyl-pyridin-2-yl)-urea; and~~
~~1-[2-(2-Chloro-pyridin-4-yl)-thiazol-4-yl]-3-(6-piperidin-1-ylmethyl-pyridin-2-yl)-urea;~~
and pharmaceutically acceptable salts thereof.

Claim 13 (canceled).

Claim 14 (currently amended): A method of inhibiting cell proliferation which comprises administering an effective amount of a compound of Formula VI



wherein R¹⁵ is one or more substituents selected from ~~H, optionally substituted heterocyclyl, phenyl, C₁-C₃ alkyl, C₁-C₂ haloalkyl, C₁-C₄ hydroxyalkyl, amino, C₁-C₄ azidoalkyl, C₁-C₄~~

~~cyanoalkyl, C₁-C₄-aminoalkyl, halo, hydroxy, (optionally substituted heterocyclyl) C₁-C₄-alkyl, optionally substituted phenoxy C₁-C₂-alkyl, C₁-C₄-alkoxy C₁-C₄-alkyl, C₁-C₄-alkylamino C₁-C₄-alkyl, C₁-C₄-hydroxyalkylamino, amino C₁-C₄-alkoxy C₁-C₄-alkyl, optionally substituted heterocycloxy, optionally substituted heterocyclyl-C₁-C₄-alkoxy, C₁-C₄-alkylamino-C₁-C₄-alkoxy, and optionally substituted phenoxy, C₁-C₄-alkoxycarbonyl, 5-6 membered heterocyclyl C₁-C₄-alkylaminocarbonyl, 5-6 membered N-containing heterocyclylcarbonyl, C₁-C₄-alkylaminocarbonyl, C₁-C₄-alkylaminothiocarbonyl, C₁-C₄-alkylamino C₁-C₄-alkylaminocarbonyl, aminocarbonyl, 5-6 membered N-containing heterocyclyl sulfonyl C₁-C₄-alkyl, 5-6 membered N-containing heterocyclyl C₁-C₄-alkylamino, C₁-C₄-alkylamino, C₁-C₄-alkylamino C₁-C₄-alkylamino C₁-C₄-alkyl, and C₁-C₄-alkylamino C₁-C₄-alkylamino;~~

wherein R¹⁶ is selected from H, ~~heterocyclylcarbonyl~~, alkylaminocarbonyl, and
alkylaminomethyl, ~~and heterocyclylmethyl~~; and

wherein R¹⁷ is selected from halo, C₁-C₆-alkyl, cycloalkylalkynyl, cycloalkyl, ~~optionally substituted indolyl, optionally substituted indazolyl, optionally substituted phenoxy, optionally substituted heteroarylsulfonyl C₁-C₄-alkyl, thienylsulfonyl- C₁-C₄-alkyl, unsubstituted 5-membered oxygen or sulfur containing heteroaryl, thienyl, unsubstituted 6-membered nitrogen containing heterocyclyl, phenyl optionally substituted with one or two substituents selected~~

~~from halo, C₁-C₄-alkylamino, amino, nitro, C₁-C₄-alkoxy, C₁-C₂-haloalkyl, hydroxy, C₁-C₄-alkylthio, C₁-C₄-alkylcarbonylamino, (optionally substituted phenyl)sulfonylamino, cyano, C₁-C₂-haloalkoxy, 5- or 6-membered N-containing heterocyclyl, aminosulfonyl, (6-membered N-containing heterocyclyl)sulfonyl, C₁-C₂-haloalkylcarbonylaminosulfonyl and (optionally substituted phenyl)aminosulfonyl, and 6-membered nitrogen-containing heterocyclyl optionally substituted with one or more substituents independently selected from pyridyl, phenyl,~~

~~C₁-C₄-alkyl, C₁-C₂-haloalkyl, C₁-C₂ alkoxy, amino, halo, piperidinyl, morpholinyl, C₁-C₂-alkylpiperazinyl, C₁-C₂-alkylaminothiocarbonyl, N,N-di C₁-C₂-alkylamino C₁-C₄-alkylonyl, N-C₁-C₂-alkylamino C₁-C₄-alkylonyl, morpholinyl C₁-C₄-alkylonylaminocarbonyl, aminocarbonyl, C₁-C₂-haloalkylcarbonylamino, morpholinyl C₁-C₄-alkylonylamino, N,N-di C₁-C₂-alkylamino and N,N-di C₁-C₂-alkylamino C₁-C₄-alkylonylamino;~~

and pharmaceutically acceptable ~~derivatives~~ salts thereof;
~~provided only one of R¹⁵ and R¹⁶ is H.~~

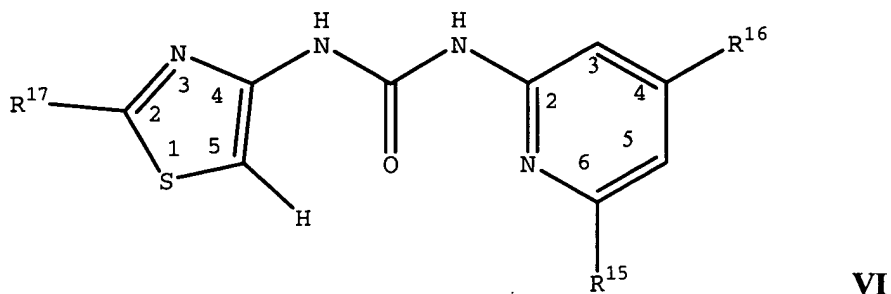
Claims 15-66 (Canceled).

Claim 67 (currently amended): The method of Claim ~~49~~ 111, wherein the compound is ~~and~~
~~pharmaceutically acceptable salts thereof~~ selected from:

~~1-pyridin-2-yl-3-(2-pyridin-4-ylthiazol-4-yl)urea;~~
~~1-(6-ethylpyridin-2-yl)-3-(2-pyridin-4-ylthiazol-4-yl)urea;~~
~~1-(2-pyridin-4-ylthiazol-4-yl)-3-(3,4,5,6-tetrahydro-2H-[1,2']bipyridinyl-6'-yl)urea;~~
~~1-(6-(diethylaminomethyl)pyridin-2-yl)-3-(2-pyridin-4-ylthiazol-4-yl)urea;~~
~~1-[6-(4-methylpiperazin-1-yl)pyridin-2-yl]-3-(2-pyridin-4-ylthiazol-4-yl)urea;~~
~~1-[6-(piperidin-1-ylmethyl)pyridin-2-yl]-3-[2-(pyridin-4-yl)thiazol-4-yl]urea;~~
1-(6-phenoxy-pyridin-2-yl)-3-(2-pyridin-4-yl-thiazol-4-yl)urea;
~~1-[2-(2-ethoxy-pyridin-4-yl)-thiazol-4-yl]-3-(6-ethyl-pyridin-2-yl)-urea;~~
~~1-(6-diethylaminomethyl-pyridin-2-yl)-3-(2-pyridin-3-yl-thiazol-4-yl)-urea;~~
~~1-[2-(2-methoxy-pyridin-4-yl)-thiazol-4-yl]-3-(6-morpholin-4-ylmethyl-pyridin-2-yl)-urea;~~
~~1-(2-pyridin-4-yl-thiazol-4-yl)-3-(6-pyrrolidin-1-ylmethyl-pyridin-2-yl)-urea;~~
~~1-(2-phenylthiazol-4-yl)-3-(6-piperidin-1-ylmethyl-pyridin-2-yl)urea;~~
1-[6-(1-methylpyrrolidin-2-ylmethoxy)pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)urea; and
~~1-[2-(4-aminophenyl)thiazol-4-yl]-3-(6-piperidin-1-ylmethyl-pyridin-2-yl)urea; and~~
1-{6-[4-(2-aminoethyl)phenoxy]pyridin-2-yl}-3-(2-pyridin-4-yl-thiazol-4-yl)urea, and
pharmaceutically acceptable salts thereof.

Claims 68-110 (canceled)

Claim 111 (currently amended): A method of treating cancer ~~inhibiting cell proliferation~~ which
comprises administering an effective amount of a compound of Formula VI



wherein R¹⁵ is one or more substituents selected from H, ~~optionally substituted heterocyclyl, phenyl, C₁-C₂-alkyl, C₁-C₂-haloalkyl, C₁-C₄-hydroxyalkyl, amino, C₁-C₄-azidoalkyl, C₁-C₄-cyanoalkyl, C₁-C₄-aminoalkyl, halo, hydroxy, (optionally substituted heterocyclyl) C₁-C₄-alkyl, optionally substituted phenoxy C₁-C₂-alkyl, C₁-C₄-alkoxy C₁-C₄-alkyl, C₁-C₄-alkylamino C₁-C₄-alkyl, C₁-C₄-hydroxyalkylamino, amino C₁-C₄-alkoxy C₁-C₄-alkyl, optionally substituted heterocycloxy, optionally substituted heterocyclyl-C₁-C₄-alkoxy, C₁-C₄-alkylamino-C₁-C₄-alkoxy, and optionally substituted phenoxy, C₁-C₄-alkoxycarbonyl, 5-6 membered heterocyclyl C₁-C₄-alkylaminocarbonyl, 5-6 membered N-containing heterocyclylcarbonyl, C₁-C₄-alkylaminocarbonyl, C₁-C₄-alkylaminethiocarbonyl, C₁-C₄-alkylamino C₁-C₄-alkylaminocarbonyl, aminocarbonyl, 5-6 membered N-containing heterocyclyl sulfonyl C₁-C₄-alkyl, 5-6 membered N-containing heterocyclyl C₁-C₄-alkylamino, C₁-C₄-alkylamino, C₁-C₄-alkylamino C₁-C₄-alkylamino C₁-C₄-alkyl, and C₁-C₄-alkylamino C₁-C₄-alkylamino;~~

wherein R¹⁶ is selected from H, ~~heterocyclylcarbonyl, alkylaminocarbonyl, and~~ alkylaminomethyl, ~~and heterocyclylmethyl;~~ and

wherein R¹⁷ is selected from halo, C₁-C₆-alkyl, cycloalkylalkynyl, cycloalkyl, ~~optionally substituted indolyl, optionally substituted indazolyl, optionally substituted phenoxy, optionally substituted heteroaryl sulfonyl C₁-C₄-alkyl, thienylsulfonyl- C₁-C₄-alkyl, unsubstituted 5-membered oxygen or sulfur-containing heteroaryl, thienyl, unsubstituted 6-membered nitrogen-containing heterocyclyl, phenyl optionally substituted with one or two substituents selected~~

~~from halo, C₁-C₄-alkylamino, amino, nitro, C₁-C₄-alkoxy, C₁-C₂-haloalkyl, hydroxy, C₁-C₄-alkylthio, C₁-C₄-alkylcarbonylamino, (optionally substituted phenyl)sulfonylamino, cyano, C₁-C₂-haloalkoxy, 5- or 6-membered N-containing~~

~~heterocyclyl, aminosulfonyl, (6-membered N-containing heterocyclyl)sulfonyl, C₁-C₂-haloalkylcarbonylamino~~sulfonyl and (optionally substituted phenyl)aminosulfonyl, and 6-membered nitrogen-containing heterocyclyl optionally substituted with ~~one or more substituents independently selected from pyridyl, phenyl,~~

~~C₁-C₄-alkyl, C₁-C₂-haloalkyl, C₁-C₂ alkoxy, amino, halo, piperidinyl, morpholinyl, C₁-C₂-alkylpiperazinyl, C₁-C₂-alkylaminothiocarbonyl, N,N-di-C₁-C₂-alkylamino-C₁-C₄-alkenyl, N-C₁-C₂-alkylamino-C₁-C₄-alkenyl, morpholinyl-C₁-C₄-alkenylaminocarbonyl, aminocarbonyl, C₁-C₂-haloalkylcarbonylamino, morpholinyl-C₁-C₄-alkenylamino, N,N-di-C₁-C₂-alkylamino and N,N-di-C₁-C₂-alkylamino-C₁-C₄-alkenylamino;~~

and pharmaceutically acceptable derivatives salts thereof;

~~provided only one of R¹⁵ and R¹⁶ is H.~~

Claim 112 (currently amended): The method of Claim 111, wherein R¹⁵ is selected from H, ~~optionally substituted pyrrolidinyl, optionally substituted piperazinyl, optionally substituted piperidinyl, morpholinyl, 1,2,3,6-tetrahydro-pyridinyl, (optionally substituted pyrrolidinyl)-C₁-C₂-alkyl, (optionally substituted piperidinyl)-C₁-C₂-alkyl, (optionally substituted piperazinyl)-C₁-C₂-alkyl, morpholinyl-C₁-C₂-alkyl, C₁-C₄-alkylamino-C₁-C₄-alkyl, C₁-C₄-hydroxyalkylamino, (optionally substituted pyrrolidinyl)-C₁-C₂-alkylamino, (optionally substituted piperidinyl)-C₁-C₂-alkylamino, (optionally substituted piperazinyl)-C₁-C₂-alkylamino, morpholinyl-C₁-C₂-alkylamino,~~ optionally substituted pyrrolidinyl-C₁-C₄-alkoxy, optionally substituted azetidiny-C₁-C₄-alkoxy, tetrahydrofuryl-C₁-C₄-alkoxy, optionally substituted piperidinyl-C₁-C₄-alkoxy, C₁-C₄-alkylamino-C₁-C₄-alkoxy, tetrahydrofuryloxy, optionally substituted piperidinyl-oxy, and optionally substituted phenoxy, ~~C₁-C₄-alkylaminocarbonyl and C₁-C₄-alkylaminothiocarbonyl;~~ wherein R¹⁶ is selected from H, ~~5-6-membered nitrogen-containing heterocyclylcarbonyl, C₁-C₄-alkylaminocarbonyl, and C₁-C₄-alkylaminomethyl, and 5-6-membered nitrogen-containing heterocyclylmethyl;~~ and wherein R¹⁷ is selected from halo, C₁-C₂-alkyl, ~~optionally substituted 5-6-membered heteroaryl-sulfonyl-C₁-C₂-alkyl, thienylsulfonyl-C₁-C₂-alkyl, optionally substituted phenoxy,~~ and C₃-C₆-cycloalkyl-C₂-C₄-alkynyl, and pharmaceutically acceptable derivatives thereof.

Claim 113 (currently amended): The method of Claim 112, wherein R¹⁵ is selected from H, tetrahydro-furanyloxy, 1-methylpyrrolidin-2-ylmethoxy, 2-pyrrolidinylmethoxy, 3-pyrrolidinylmethoxy, 1-Boc-pyrrolidin-2-ylmethoxy, 4-piperidinylmethoxy, 1-Boc-piperidin-4-ylmethoxy, 1-Boc-piperidin-4-ylethoxy, piperidin-4-ylethoxy, 1-methyl-piperidin-4-ylmethoxy, 1-Boc-azetidin-3-ylmethoxy, 1-methyl-azetidin-3-ylmethoxy, 3-azetidinylmethoxy, 1-methyl-piperidin-4-yloxy, ~~phenyloxy, phenoxy, 4-(pyrrolidin-1-ylmethyl)phenoxy, and~~ dimethylaminoethoxy, ~~1-piperidinylmethyl, 1-(piperidin-1-yl)ethyl, 3-methylpiperidin-1-ylmethyl, 1-pyrrolidinylmethyl, 2,2,6,6-tetramethylpiperidin-1-ylmethyl, 2,6-dimethylpiperidin-1-ylmethyl, dimethylaminomethyl, diethylaminomethyl, diethylaminothiocarbonyl, diethylaminocarbonyl, N-Boc-N-isopropylaminomethyl, isopropylaminomethyl, 2-thienylsulfonylmethyl, hydroxypropylamino, 4-ethyl-piperidin-1-yl, 4-(2-pyridyl)piperidin-1-yl, 1-methylpiperidin-4-yl, 4-(2-pyrazinyl)piperidin-1-yl, 1-methyl-1,2,3,6-tetrahydro-pyridin-4-yl, 1,2,3,6-tetrahydro-pyridin-4-yl, and 1-Boc-1,2,3,6-tetrahydro-pyridin-4-yl;~~ wherein R¹⁶ is selected from H, ~~1-piperidinylecarbonyl, diethylaminocarbonyl, and~~ diethylaminomethyl, ~~1-piperidinylmethyl;~~ and wherein R¹⁷ is selected from chloro, bromo, methyl and cyclopropylethynyl, ~~and pharmaceutically acceptable derivatives thereof.~~

Claim 114 (currently amended): The method of Claim 113, wherein R¹⁷ is chloro or bromo, ~~and pharmaceutically acceptable derivatives thereof.~~

Claim 115 (currently amended): The method of Claim 111, wherein R¹⁵ is selected from H, ~~optionally substituted pyrrolidinyl, optionally substituted piperazinyl, optionally substituted piperidinyl, morpholinyl, 1,2,3,6-tetrahydro-pyridinyl, (optionally substituted pyrrolidinyl)-C₁-C₂-alkyl, (optionally substituted piperidinyl)-C₁-C₂-alkyl, (optionally substituted piperazinyl)-C₁-C₂-alkyl, morpholinyl-C₁-C₂-alkyl, (optionally substituted pyrrolidinyl)-C₁-C₂-alkylamino, (optionally substituted piperidinyl)-C₁-C₂-alkylamino, (optionally substituted piperazinyl)-C₁-C₂-alkylamino, morpholinyl-C₁-C₂-alkylamino, C₁-C₄-alkylamino-C₁-C₄-alkyl, C₁-C₄-hydroxyalkylamino, optionally substituted pyrrolidinyl-C₁-C₄-alkoxy, optionally substituted azetidiny-C₁-C₄-alkoxy, tetrahydrofuryl-C₁-C₄-alkoxy, optionally substituted piperidinyl-C₁-C₄-alkoxy, C₁-C₄-alkylamino-C₁-C₄-alkoxy, tetrahydrofuryloxy, optionally substituted piperidin-yloxy, optionally substituted phenoxy, ~~C₁-C₄-alkylaminocarbonyl and C₁-C₄-alkylaminothiocarbonyl;~~ wherein R¹⁶ is selected from H, ~~5-6-membered nitrogen-containing~~~~

~~heterocyclylcarbonyl, C₁-C₄-alkylaminocarbonyl, and C₁-C₄-alkylaminomethyl, and 5-6~~
~~membered nitrogen-containing heterocyclylmethyl; and wherein R¹⁷ is selected from C₃-C₆-~~
~~cycloalkyl and phenyl optionally substituted with one or two substituents selected from halo,~~
~~C₁-C₄-alkylamino, amino, nitro, C₁-C₄-alkoxy, C₁-C₂-haloalkyl, hydroxy, C₁-C₄-~~
~~alkylthio, C₁-C₄-alkylcarbonylamino, (optionally substituted phenyl)sulfonylamino,~~
~~cyano, C₁-C₂-haloalkoxy, 5- or 6-membered N-containing heterocyclyl, aminosulfonyl,~~
~~(6-membered N-containing heterocyclyl)sulfonyl, C₁-C₂-haloalkylcarbonylaminosulfonyl~~
~~and (optionally substituted phenyl)aminosulfonyl;~~
~~and pharmaceutically acceptable derivatives thereof.~~

Claim 116 (currently amended): The method of Claim 115, wherein R¹⁵ is selected from H,
tetrahydro-furanyloxy, 1-methylpyrrolidin-2-ylmethoxy, 2-pyrrolidinylmethoxy, 3-
pyrrolidinylmethoxy, 1-Boc-pyrrolidin-2-ylmethoxy, 4-piperidinylmethoxy, 1-Boc-piperidin-4-
ylmethoxy, 1-Boc-piperidin-4-ylethoxy, piperidin-4-ylethoxy, 1-methyl-piperidin-4-ylmethoxy,
1-Boc-azetidin-3-ylmethoxy, 1-methyl-azetidin-3-ylmethoxy, 3-azetidinylmethoxy, 1-methyl-
piperidin-4-yloxy, ~~phenyloxy, phenoxy, 4-(pyrrolidin-1-ylmethyl)phenoxy, and~~
~~dimethylaminoethoxy, 1-piperidinylmethyl, 1-(piperidin-1-yl)ethyl, 3-methylpiperidin-1-~~
~~ylmethyl, 1-pyrrolidinylmethyl, 2,2,6,6-tetramethylpiperidin-1-ylmethyl, 2,6-dimethylpiperidin-~~
~~1-ylmethyl, dimethylaminomethyl, diethylaminomethyl, diethylaminothiocarbonyl,~~
~~diethylaminocarbonyl, N-Boc-N-isopropylaminomethyl, isopropylaminomethyl, 2-~~
~~thienylsulfonylmethyl, hydroxypropylamine, 4-ethyl-piperidin-1-yl, 4-(2-pyridyl)piperidin-1-yl,~~
~~1-methylpiperidin-4-yl, 4-(2-pyrazinyl)piperidin-1-yl, 1-methyl-1,2,3,6-tetrahydro-pyridin-4-yl,~~
~~1,2,3,6-tetrahydro-pyridin-4-yl, and 1-Boc-1,2,3,6-tetrahydro-pyridin-4-yl; wherein R¹⁶ is~~
selected from H, ~~1-piperidinylcarbonyl, diethylaminocarbonyl, and diethylaminomethyl, 1-~~
~~piperidinylmethyl;~~ and wherein R¹⁷ is selected from cyclopropyl and phenyl ~~optionally~~
~~substituted with aminosulfonyl, and pharmaceutically acceptable derivatives thereof.~~

Claim 117 (currently amended): The method of Claim 116, wherein R¹⁷ is unsubstituted phenyl;
~~and pharmaceutically acceptable derivatives thereof.~~

Claim 118 (currently amended): The method of Claim 111, wherein R¹⁵ is selected from H,
~~optionally substituted pyrrolidinyl, optionally substituted piperazinyl, optionally substituted~~

~~piperidinyl, morpholinyl, 1,2,3,6-tetrahydro-pyridinyl, (optionally substituted pyrrolidinyl) C₁-C₂-alkyl, (optionally substituted piperidinyl) C₁-C₂-alkyl, (optionally substituted piperazinyl) C₁-C₂-alkyl, morpholinyl C₁-C₂-alkyl, (optionally substituted pyrrolidinyl) C₁-C₂-alkylamino, (optionally substituted piperidinyl) C₁-C₂-alkylamino, (optionally substituted piperazinyl) C₁-C₂-alkylamino, morpholinyl C₁-C₂-alkylamino, C₁-C₄-alkylamino C₁-C₄-alkyl, C₁-C₄-hydroxyalkylamino, optionally substituted pyrrolidinyl-C₁-C₄-alkoxy, optionally substituted azetidiny-C₁-C₄-alkoxy, tetrahydrofuryl-C₁-C₄-alkoxy, optionally substituted piperidinyl-C₁-C₄-alkoxy, C₁-C₄-alkylamino-C₁-C₄-alkoxy, tetrahydrofuryloxy, optionally substituted piperidinyl, and optionally substituted phenoxy, C₁-C₄-alkylaminocarbonyl and C₁-C₄-alkylaminethiocarbonyl; wherein R¹⁶ is selected from H, 5-6-membered nitrogen-containing heterocyclylcarbonyl, C₁-C₄-alkylaminocarbonyl, and C₁-C₄-alkylaminomethyl, and 5-6-membered nitrogen-containing heterocyclylmethyl; and wherein R¹⁷ is selected from optionally substituted indazolyl, optionally substituted indolyl, unsubstituted 5-membered oxygen or sulfur containing heterocyclyl, unsubstituted thienyl, unsubstituted 6-membered nitrogen-containing heterocyclyl, and 6-membered nitrogen-containing heterocyclyl substituted with one or more substituents independently selected from pyridyl, phenyl, C₁-C₄-alkyl, C₁-C₂-haloalkyl, C₁-C₂ alkoxy, amino, halo, piperidinyl, morpholinyl, C₁-C₂-alkylpiperazinyl, C₁-C₂-alkylaminethiocarbonyl, N,N-di-C₁-C₂-alkylamino C₁-C₄-alkylonyl, N-C₁-C₂-alkylamino C₁-C₄-alkylonyl, morpholinyl C₁-C₄-alkylonylaminocarbonyl, aminocarbonyl, C₁-C₂-haloalkylcarbonylamino, morpholinyl C₁-C₄-alkylonylamino, N,N-di-C₁-C₂-alkylamino and N,N-di-C₁-C₂-alkylamino C₁-C₄-alkylonylamino; and pharmaceutically acceptable derivatives thereof.~~

Claim 119 (currently amended): The method of Claim 118, wherein R¹⁵ is selected from H, tetrahydro-furanyloxy, 1-methylpyrrolidin-2-ylmethoxy, 2-pyrrolidinylmethoxy, 3-pyrrolidinylmethoxy, 1-Boc-pyrrolidin-2-ylmethoxy, 4-piperidinylmethoxy, 1-Boc-piperidin-4-ylmethoxy, 1-Boc-piperidin-4-ylethoxy, piperidin-4-ylethoxy, 1-methyl-piperidin-4-ylmethoxy, 1-Boc-azetidin-3-ylmethoxy, 1-methyl-azetidin-3-ylmethoxy, 3-azetidylmethoxy, 1-methyl-piperidin-4-yloxy, phenyloxy, 4-(pyrrolidin-1-ylmethyl)phenoxy, and dimethylaminoethoxy, ~~1-piperidinylmethyl, 1-(piperidin-1-yl)ethyl, 3-methylpiperidin-1-ylmethyl, 1-pyrrolidinylmethyl, 2,2,6,6-tetramethylpiperidin-1-ylmethyl, 2,6-dimethylpiperidin-1-ylmethyl,~~

~~dimethylaminomethyl, diethylaminomethyl, diethylaminothiocarbonyl, diethylaminocarbonyl, N-Boc-N-isopropylaminomethyl, isopropylaminomethyl, 2-thienylsulfonylmethyl, hydroxypropylamino, 4-ethyl piperidin-1-yl, 4-(2-pyridyl)piperidin-1-yl, 1-methylpiperidin-4-yl, 4-(2-pyrazinyl)piperidin-1-yl, 1-methyl-1,2,3,6-tetrahydro-pyridin-4-yl, 1,2,3,6-tetrahydro-pyridin-4-yl, and 1-Boc-1,2,3,6-tetrahydro-pyridin-4-yl; wherein R¹⁶ is selected from H, 4-piperidinylcarbonyl, diethylaminocarbonyl, and diethylaminomethyl, 1-piperidinylmethyl; and wherein R¹⁷ is selected from 5-indazolyl, 1-Boc-indol-5-yl, unsubstituted thienyl, 5-tert-butylloxazol-2-yl and 4-pyridyl substituted with one or more substituents independently selected from methoxy and chloro; and pharmaceutically acceptable derivatives thereof.~~

Claim 120 (currently amended): The method of Claim ~~119~~ 111, wherein R¹⁷ is 4-pyridyl, ~~and pharmaceutically acceptable derivatives thereof.~~

Claim 121 (currently amended): The method of Claim 111, wherein the compound is ~~and pharmaceutically acceptable derivatives thereof~~ selected from:

~~1-[6-(3-Methyl-piperidin-1-ylmethyl)-pyridin-2-yl]-3-(2-phenyl-thiazol-4-yl)-urea;
1-[4-(Piperidine-1-carbonyl)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;
1-(2-Chloro-thiazol-4-yl)-3-[4-(piperidine-1-carbonyl)-pyridin-2-yl]-urea;
N,N-Diethyl-2-[3-(2-pyridin-4-yl-thiazol-4-yl)-ureido]-isonicotinamide;
N,N-Diethyl-2-[3-(2-phenyl-thiazol-4-yl)-ureido]-isonicotinamide;
2-[3-(2-Bromo-thiazol-4-yl)-ureido]-N,N-diethyl-isonicotinamide;
1-(4-Diethylaminomethyl-pyridin-2-yl)-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;
1-[6-(2,6-Dimethyl-piperidin-1-ylmethyl)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;
1-[6-(1-Piperidin-1-yl-ethyl)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;
2-((6-[3-(2-Pyridin-4-yl-thiazol-4-yl)-ureido]-pyridin-2-ylamino)-methyl)-piperidine-1-carboxylic acid tert-butyl ester;
1-[6-[(Piperidin-2-ylmethyl)-amino]-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;
(S)-1-[6-(3-Methyl-piperidin-1-ylmethyl)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;
(R)-1-[6-(3-Methyl-piperidin-1-ylmethyl)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;
1-(2-Chloro-thiazol-4-yl)-3-(6-piperidin-1-ylmethyl-pyridin-2-yl)-urea;~~

1-(2-Bromo-thiazol-4-yl)-3-[6-(2-piperidin-4-yl-ethoxy)-pyridin-2-yl]-urea;
1-(2-Chloro-thiazol-4-yl)-3-[6-(2-piperidin-4-yl-ethoxy)-pyridin-2-yl]-urea;
1-[6-(Azetidin-3-ylmethoxy)-pyridin-2-yl]-3-(2-bromo-thiazol-4-yl)-urea;
1-[6-(Azetidin-3-ylmethoxy)-pyridin-2-yl]-3-(2-chloro-thiazol-4-yl)-urea;
1-(2-Bromo-thiazol-4-yl)-3-[6-(piperidin-4-ylmethoxy)-pyridin-2-yl]-urea;
1-(2-Chloro-thiazol-4-yl)-3-[6-(piperidin-4-ylmethoxy)-pyridin-2-yl]-urea;
tert-Butyl 3-{6-[3-(2-pyridin-4-yl-thiazol-4-yl)-ureido]-pyridin-2-yloxymethyl}-pyrrolidine-1-carboxylate;
1-(2-Pyridin-4-yl-thiazol-4-yl)-3-[6-(pyrrolidin-3-ylmethoxy)-pyridin-2-yl]-urea;
1-(2-Cyclopropyl-thiazol-4-yl)-3-[6-(2-piperidin-4-yl-ethoxy)-pyridin-2-yl]-urea;
~~1-[6-(Isopropylamino-methyl)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;~~
~~1-[6-(Isopropylamino-methyl)-pyridin-2-yl]-3-(2-phenyl-thiazol-4-yl)-urea;~~
~~1-(2-Bromo-thiazol-4-yl)-3-[6-(isopropylamino-methyl)-pyridin-2-yl]-urea;~~
1-(2-Bromo-thiazol-4-yl)-3-[6-(1-methyl-pyrrolidin-2-ylmethoxy)-pyridin-2-yl]-urea;
1-(2-Chloro-thiazol-4-yl)-3-[6-(1-methyl-pyrrolidin-2-ylmethoxy)-pyridin-2-yl]-urea;
1-(2-phenylthiazol-4-yl)-3-(6-*p*-pyrrolidin-1-ylmethylphenoxy)pyridin-2-yl)urea;
1-(2-Pyridin-4-yl-thiazol-4-yl)-3-[6-(tetrahydro-furan-3-yloxy)-pyridin-2-yl]-urea;
~~1-[2-(1H-Indazol-5-yl)-thiazol-4-yl]-3-[6-(piperidin-1-ylmethyl)-pyridin-2-yl]-urea;~~
~~1-(1'-Methyl-1',2',3',6'-tetrahydro-[2,4']bipyridinyl-6-yl)-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;~~
~~1-(2-Bromo-thiazol-4-yl)-3-(1'-methyl-1',2',3',6'-tetrahydro-[2,4']bipyridinyl-6-yl)-urea;~~
~~1-(1'-Methyl-1',2',3',6'-tetrahydro-2[2,4]bipyridinyl-6-yl)-3-(2-phenyl-thiazol-4-yl)-urea;~~
~~1-[6-(3-Hydroxy-propylamino)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;~~
~~1-(2-Bromo-thiazol-4-yl)-3-[6-(3-hydroxy-propylamino)-pyridin-2-yl]-urea;~~
~~1-(1'-Methyl-1',2',3',4',5',6'-hexahydro-[2,4']bipyridinyl-6-yl)-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;~~
~~urea;~~
~~1-(1'-Methyl-1',2',3',4',5',6'-hexahydro-[2,4']bipyridinyl-6-yl)-3-(2-phenyl-thiazol-4-yl)-urea;~~
~~6-[3-(2-Pyridin-4-yl-thiazol-4-yl)-ureido]-3',6'-dihydro-2'H-[2,4]bipyridinyl-1'-carboxylic acid~~
~~*tert*-butylester;~~
~~1-(2-Pyridin-4-yl-thiazol-4-yl)-3-(1',2',3',6'-tetrahydro-[2,4']bipyridinyl-6-yl)-urea;~~
1-(2-Pyridin-4-yl-thiazol-4-yl)-3-[6-(tetrahydro-furan-3-ylmethoxy)-pyridin-2-yl]-urea;

2-[6-[3-(2-Pyridin-4-yl-thiazol-4-yl)-ureido]-pyridin-2-yloxymethyl]-pyrrolidine-1-carboxylic acid tert-butyl ester;

1-(2-Pyridin-4-yl-thiazol-4-yl)-3-[6-(pyrrolidin-2-ylmethoxy)-pyridin-2-yl]-urea;

~~6-[3-(2-Pyridin-4-yl-thiazol-4-yl)-ureido]-pyridine-2-carboethioic acid diethylamide;~~

~~1-(2-Bromo-thiazol-4-yl)-3-[6-(3-methyl-piperidin-1-ylmethyl)-pyridin-2-yl]-urea;~~

~~1-(2-Chloro-thiazol-4-yl)-3-[6-(3-methyl-piperidin-1-ylmethyl)-pyridin-2-yl]-urea;~~

~~1-(2-Phenyl-thiazol-4-yl)-3-[4-(piperidine-1-carbonyl)-pyridin-2-yl]-urea;~~

~~1-(2-Bromo-thiazol-4-yl)-3-[4-(piperidine-1-carbonyl)-pyridin-2-yl]-urea;~~

1-[2-(2-Methoxy-pyridin-4-yl)-thiazol-4-yl]-3-(6-phenoxy-pyridin-2-yl)-urea;

1-[2-(2-Methoxy-pyridin-4-yl)-thiazol-4-yl]-3-[6-(1-methyl-pyrrolidin-2-ylmethoxy)-pyridin-2-yl]-urea;

1-[6-(2-Dimethylamino-ethoxy)-pyridin-2-yl]-3-[2-(2-methoxy-pyridin-4-yl)-thiazol-4-yl]-urea;

1-[6-(1-Methyl-pyrrolidin-2-ylmethoxy)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;

~~1-(2-phenylthiazol-4-yl)-3-(6-pyrrolidin-1-ylmethyl-pyridin-2-yl)urea;~~

~~1-(6-Diethylaminomethylpyridin-2-yl)-3-(2-phenylthiazol-4-yl)urea;~~

(S)-1-[6-(1-Methylpyrrolidin-2-ylmethoxy)pyridin-2-yl]-3-(2-phenylthiazol-4-yl)urea;

1-[6-(2-Piperidin-4-yl-ethoxy)pyridin-2-yl]-3-[2-phenylthiazol-4-yl]urea;

~~1-[6-(4-Ethylpiperazin-1-yl)-pyridin-2-yl]-3-(2-phenylthiazol-4-yl)urea;~~

~~Diethyl 6-[3-(2-phenylthiazol-4-yl)ureido]-pyridine-2-carboxamide;~~

1-(2-Pyridin-4-yl-thiazol-4-yl)-3-(6-*p*-pyrrolidin-1-ylmethylphenoxy)pyridin-2-yl)urea;

1-(2-Bromothiazol-4-yl)-3-(6-*p*-pyrrolidin-1-ylmethylphenoxy)pyridin-2-yl)urea;

1-[6-(Piperidin-4-ylmethoxy)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;

1-[6-(1-Methyl-piperidin-4-ylmethoxy)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;

1-[6-(1-Methyl-piperidin-4-yloxy)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;

1-[6-(1-Methyl-azetidin-3-ylmethoxy)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;

1-[6-(Azetidin-3-ylmethoxy)-pyridin-2-yl]-3-(2-phenyl-thiazol-4-yl)-urea;

1-[6-(1-Methyl-azetidin-3-ylmethoxy)-pyridin-2-yl]-3-(2-phenyl-thiazol-4-yl)-urea;

1-(2-Phenyl-thiazol-4-yl)-3-[6-(piperidin-4-ylmethoxy)-pyridin-2-yl]-urea;

1-[6-(1-Methyl-piperidin-4-ylmethoxy)-pyridin-2-yl]-3-(2-phenyl-thiazol-4-yl)-urea;

1-[6-(1-Methyl-piperidin-4-yloxy)-pyridin-2-yl]-3-(2-phenyl-thiazol-4-yl)-urea;

1-[6-(2-Piperidin-4-yl-ethoxy)-pyridin-2-yl]-3-(2-thiophen-2-yl-thiazol-4-yl)-urea; and

1-[6-(1-Methyl-pyrrolidin-2-ylmethoxy)-pyridin-2-yl]-3-[2-(thiophene-2-sulfonylmethyl)-thiazol-4-yl]-urea;

~~1-[2-(2-Methoxy-pyridin-4-yl)-thiazol-4-yl]-3-[6-(piperidin-1-ylmethyl-pyridin-2-yl)-urea]; and~~
~~[2-(2-Chloro-pyridin-4-yl)-thiazol-4-yl]-3-[6-(piperidin-1-ylmethyl-pyridin-2-yl)-urea~~
pharmaceutically acceptable salts thereof.

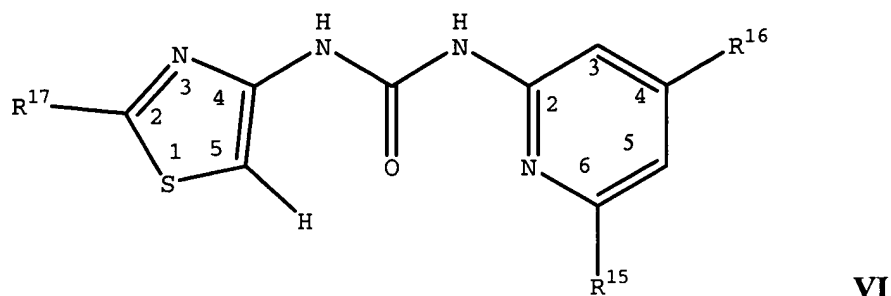
Claim 122 (currently amended): The method of Claim 111, wherein the compound is and
~~pharmaceutically acceptable derivatives thereof~~ selected from:

~~1-[6-(3-Methyl-piperidin-1-ylmethyl)-pyridin-2-yl]-3-(2-phenyl-thiazol-4-yl)-urea;~~
~~1-[4-(Piperidine-1-carbonyl)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;~~
~~N,N-Diethyl-2-[3-(2-pyridin-4-yl-thiazol-4-yl)-ureido]-isonicotinamide;~~
~~1-(4-Diethylaminomethyl-pyridin-2-yl)-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;~~
~~1-[6-(2,6-Dimethyl-piperidin-1-ylmethyl)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;~~
~~1-[6-(1-Piperidin-1-yl-ethyl)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;~~
~~2-[(6-[3-(2-Pyridin-4-yl-thiazol-4-yl)-ureido]-pyridin-2-ylamino)-methyl]-piperidine-1-carboxylic acid tert-butyl ester;~~
~~1-{6-[(Piperidin-2-ylmethyl)-amino]-pyridin-2-yl}-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;~~
~~(S)-1-[6-(3-Methyl-piperidin-1-ylmethyl)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;~~
~~(R)-1-[6-(3-Methyl-piperidin-1-ylmethyl)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;~~
~~1-(2-Chloro-thiazol-4-yl)-3-[6-(piperidin-1-ylmethyl-pyridin-2-yl)-urea];~~
1-(2-Bromo-thiazol-4-yl)-3-[6-(2-piperidin-4-yl-ethoxy)-pyridin-2-yl]-urea;
1-(2-Chloro-thiazol-4-yl)-3-[6-(2-piperidin-4-yl-ethoxy)-pyridin-2-yl]-urea;
1-[6-(Azetidin-3-ylmethoxy)-pyridin-2-yl]-3-(2-bromo-thiazol-4-yl)-urea;
1-[6-(Azetidin-3-ylmethoxy)-pyridin-2-yl]-3-(2-chloro-thiazol-4-yl)-urea;
1-(2-Bromo-thiazol-4-yl)-3-[6-(piperidin-4-ylmethoxy)-pyridin-2-yl]-urea;
1-(2-Chloro-thiazol-4-yl)-3-[6-(piperidin-4-ylmethoxy)-pyridin-2-yl]-urea;
3-(4-{3-[6-(1-Methyl-pyrrolidin-2-ylmethoxy)-pyridin-2-yl]-ureido}-thiazol-2-yl)-
benzenesulfonamide;
tert-Butyl 3-{6-[3-(2-pyridin-4-yl-thiazol-4-yl)-ureido]-pyridin-2-yloxymethyl}-pyrrolidine-1-carboxylate;
1-(2-Pyridin-4-yl-thiazol-4-yl)-3-[6-(pyrrolidin-3-ylmethoxy)-pyridin-2-yl]-urea;

1-(2-Cyclopropyl-thiazol-4-yl)-3-[6-(2-piperidin-4-yl-ethoxy)-pyridin-2-yl]-urea;
~~Isopropyl {6 [3 (2 pyridin 4 yl thiazol 4 yl) ureido] pyridin 2 ylmethyl} carbamic acid tert-butyl ester;~~
~~1-[6 (Isopropylamine methyl) pyridin 2 yl] 3 (2 pyridin 4 yl thiazol 4 yl) urea;~~
~~Isopropyl {6 [3 (2 phenyl thiazol 4 yl) ureido] pyridin 2 ylmethyl} carbamic acid tert-butyl ester;~~
~~1-[6 (Isopropylamine methyl) pyridin 2 yl] 3 (2 phenyl thiazol 4 yl) urea;~~
1-(2-Bromo-thiazol-4-yl)-3-[6-(1-methyl-pyrrolidin-2-ylmethoxy)-pyridin-2-yl]-urea;
1-(2-Chloro-thiazol-4-yl)-3-[6-(1-methyl-pyrrolidin-2-ylmethoxy)-pyridin-2-yl]-urea;
1-(2-phenylthiazol-4-yl)-3-(6-*p*-pyrrolidin-1-ylmethylphenoxypyridin-2-yl)urea;
1-(2-Pyridin-4-yl-thiazol-4-yl)-3-[6-(tetrahydro-furan-3-yloxy)-pyridin-2-yl]-urea;
~~1-[2 (1H Indazol 5 yl) thiazol 4 yl] 3 (6 piperidin 1 ylmethyl pyridin 2 yl) urea;~~
~~1-(1' Methyl 1',2',3',6' tetrahydro [2,4']bipyridinyl 6 yl) 3 (2 pyridin 4 yl thiazol 4 yl) urea;~~
~~1-(2 Bromo thiazol 4 yl) 3 (1' methyl 1',2',3',6' tetrahydro [2,4']bipyridinyl 6 yl) urea;~~
~~1-(1' Methyl 1',2',3',6' tetrahydro 2[2,4]bipyridinyl 6 yl) 3 (2 phenyl thiazol 4 yl) urea;~~
~~1-[6 (3 Hydroxy propylamine) pyridin 2 yl] 3 (2 pyridin 4 yl thiazol 4 yl) urea;~~
~~1-(2 Bromo thiazol 4 yl) 3 [6(3 hydroxy propylamine) pyridin 2 yl] urea;~~
~~1-(1' Methyl 1',2',3',4',5',6' hexahydro [2,4']bipyridinyl 6 yl) 3 (2 pyridin 4 yl thiazol 4 yl) urea;~~
~~1-(1' Methyl 1',2',3',4',5',6' hexahydro [2,4']bipyridinyl 6 yl) 3 (2 phenyl thiazol 4 yl) urea;~~
~~6 [3 (2 Pyridin 4 yl thiazol 4 yl) ureido] 3',6' dihydro 2'H [2,4]bipyridinyl 1' carboxylic acid tert-butylester;~~
~~1-(2 Pyridin 4 yl thiazol 4 yl) 3 (1',2',3',6' tetrahydro [2,4']bipyridinyl 6 yl) urea;~~
1-(2-Pyridin-4-yl-thiazol-4-yl)-3-[6-(tetrahydro-furan-3-ylmethoxy)-pyridin-2-yl]-urea;
2-[6-[3-(2-Pyridin-4-yl-thiazol-4-yl)-ureido]-pyridin-2-yloxymethyl]-pyrrolidine-1-carboxylic acid tert-butyl ester;
1-(2-Pyridin-4-yl-thiazol-4-yl)-3-[6-(pyrrolidin-2-ylmethoxy)-pyridin-2-yl]-urea;
~~6 [3 (2 Pyridin 4 yl thiazol 4 yl) ureido] pyridine 2 carbothioic acid diethylamide;~~
~~1-(2 Bromo thiazol 4 yl) 3 [6 (3 methyl piperidin 1 ylmethyl) pyridin 2 yl] urea;~~
~~1-(2 Chloro thiazol 4 yl) 3 [6 (3 methyl piperidin 1 ylmethyl) pyridin 2 yl] urea;~~
1-[2-(2-Methoxy-pyridin-4-yl)-thiazol-4-yl]-3-[6-(1-methyl-pyrrolidin-2-ylmethoxy)-pyridin-2-yl]-urea;

1-[6-(2-Dimethylamino-ethoxy)-pyridin-2-yl]-3-[2-(2-methoxy-pyridin-4-yl)-thiazol-4-yl]-urea;
1-[6-(1-Methyl-pyrrolidin-2-ylmethoxy)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;
~~1-(2-phenylthiazol-4-yl)-3-(6-pyrrolidin-1-ylmethyl-pyridin-2-yl)urea;~~
~~1-(6-Diethylaminomethylpyridin-2-yl)-3-(2-phenylthiazol-4-yl)urea;~~
(S)-1-[6-(1-Methylpyrrolidin-2-ylmethoxy)pyridin-2-yl]-3-(2-phenylthiazol-4-yl)urea;
1-[6-(2-Piperidin-4-yl-ethoxy)pyridin-2-yl]-3-[2-phenylthiazol-4-yl]urea;
~~1-[6-(4-Ethylpiperazin-1-yl)-pyridin-2-yl]-3-(2-phenylthiazol-4-yl)urea;~~
~~1-(2-phenylthiazol-4-yl)-3-[6-(4-pyrimidin-2-yl-piperazin-1-yl)pyridin-2-yl]urea;~~
~~Diethyl 6-[3-(2-phenylthiazol-4-yl)urcide]pyridine-2-carboxamide;~~
1-(2-Pyridin-4-yl-thiazol-4-yl)-3-(6-*p*-pyrrolidin-1-ylmethylphenoxypyridin-2-yl)urea;
1-(2-Bromothiazol-4-yl)-3-(6-*p*-pyrrolidin-1-ylmethylphenoxypyridin-2-yl)urea;
1-[6-(Piperidin-4-ylmethoxy)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;
1-[6-(1-Methyl-piperidin-4-ylmethoxy)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;
1-[6-(1-Methyl-piperidin-4-yloxy)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;
1-[6-(1-Methyl-azetidin-3-ylmethoxy)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;
1-[6-(Azetidin-3-ylmethoxy)-pyridin-2-yl]-3-(2-phenyl-thiazol-4-yl)-urea;
1-[6-(1-Methyl-azetidin-3-ylmethoxy)-pyridin-2-yl]-3-(2-phenyl-thiazol-4-yl)-urea;
1-(2-Phenyl-thiazol-4-yl)-3-[6-(piperidin-4-ylmethoxy)-pyridin-2-yl]-urea;
1-[6-(1-Methyl-piperidin-4-ylmethoxy)-pyridin-2-yl]-3-(2-phenyl-thiazol-4-yl)-urea;
1-[6-(1-Methyl-piperidin-4-yloxy)-pyridin-2-yl]-3-(2-phenyl-thiazol-4-yl)-urea;
1-[6-(2-Piperidin-4-yl-ethoxy)-pyridin-2-yl]-3-(2-thiophen-2-yl-thiazol-4-yl)-urea; and
1-[6-(1-Methyl-pyrrolidin-2-ylmethoxy)-pyridin-2-yl]-3-[2-(thiophene-2-sulfonylmethyl)-thiazol-4-yl]-urea;
~~1-[2-(2-Methoxy-pyridin-4-yl)-thiazol-4-yl]-3-(6-piperidin-1-ylmethyl-pyridin-2-yl)-urea; and~~
~~[2-(2-Chloro-pyridin-4-yl)-thiazol-4-yl]-3-(6-piperidin-1-ylmethyl-pyridin-2-yl)-urea;~~
and pharmaceutically acceptable salts thereof.

Claim 123 (currently amended): A method of inhibiting a serine/threonine kinase which comprises administering an effective amount of a compound of Formula VI



wherein R¹⁵ is one or more substituents selected from ~~H, optionally substituted heterocyclyl, phenyl, C₁-C₂-alkyl, C₁-C₂-haloalkyl, C₁-C₄-hydroxyalkyl, amino, C₁-C₄-azidoalkyl, C₁-C₄-cyanoalkyl, C₁-C₄-aminoalkyl, halo, hydroxy, (optionally substituted heterocyclyl) C₁-C₄-alkyl, optionally substituted phenoxy, C₁-C₂-alkyl, C₁-C₄-alkoxy, C₁-C₄-alkylamino, C₁-C₄-alkyl, C₁-C₄-hydroxyalkylamino, amino, C₁-C₄-alkoxy, C₁-C₄-alkyl, optionally substituted heterocycloxy, optionally substituted heterocyclyl-C₁-C₄-alkoxy, C₁-C₄-alkylamino-C₁-C₄-alkoxy, and optionally substituted phenoxy, C₁-C₄-alkoxycarbonyl, 5-6 membered heterocyclyl, C₁-C₄-alkylaminocarbonyl, 5-6 membered N-containing heterocyclylcarbonyl, C₁-C₄-alkylaminocarbonyl, C₁-C₄-alkylaminethiocarbonyl, C₁-C₄-alkylamino, C₁-C₄-alkylaminocarbonyl, aminocarbonyl, 5-6 membered N-containing heterocyclyl sulfonyl, C₁-C₄-alkyl, 5-6 membered N-containing heterocyclyl, C₁-C₄-alkylamino, C₁-C₄-alkylamino, C₁-C₄-alkylamino, C₁-C₄-alkylamino, C₁-C₄-alkyl, and C₁-C₄-alkylamino, C₁-C₄-alkylamino;~~

wherein R¹⁶ is selected from H, ~~heterocyclylcarbonyl, alkylaminocarbonyl, and~~ alkylaminomethyl, ~~and heterocyclylmethyl;~~ and

wherein R¹⁷ is selected from halo, C₁-C₆-alkyl, cycloalkylalkynyl, cycloalkyl, ~~optionally substituted indolyl, optionally substituted indazolyl, optionally substituted phenoxy, optionally substituted heteroaryl, sulfonyl, C₁-C₄-alkyl, thienyl, sulfonyl, C₁-C₄-alkyl, unsubstituted 5-membered oxygen or sulfur-containing heteroaryl, thienyl, unsubstituted 6-membered nitrogen-containing heterocyclyl, phenyl optionally substituted with one or two substituents selected~~

~~from halo, C₁-C₄-alkylamino, amino, nitro, C₁-C₄-alkoxy, C₁-C₂-haloalkyl, hydroxy, C₁-C₄-alkylthio, C₁-C₄-alkylcarbonylamino, (optionally substituted phenyl)sulfonylamino, cyano, C₁-C₂-haloalkoxy, 5- or 6-membered N-containing~~

~~heterocyclyl, aminosulfonyl, (6-membered N-containing heterocyclyl)sulfonyl, C₁-C₂-haloalkylcarboxylaminosulfonyl and (optionally substituted phenyl)aminosulfonyl, and 6-membered nitrogen-containing heterocyclyl optionally substituted with one or more substituents independently selected from pyridyl, phenyl,~~

~~C₁-C₄-alkyl, C₁-C₂-haloalkyl, C₁-C₂ alkoxy, amino, halo, piperidinyl, morpholinyl, C₁-C₂-alkylpiperazinyl, C₁-C₂-alkylaminothiocarbonyl, N,N-di-C₁-C₂-alkylamino-C₁-C₄-alkenyl, N-C₁-C₂-alkylamino-C₁-C₄-alkenyl, morpholinyl-C₁-C₄-alkenylaminocarbonyl, aminocarbonyl, C₁-C₂-haloalkylcarboxylamino, morpholinyl-C₁-C₄-alkenylamino, N,N-di-C₁-C₂-alkylamino and N,N-di-C₁-C₂-alkylamino-C₁-C₄-alkenylamine;~~

and pharmaceutically acceptable derivatives salts thereof;

~~provided only one of R¹⁵ and R¹⁶ is H.~~

Claim 124 (currently amended): The method of Claim 123 wherein R¹⁵ is selected from ~~H,~~
~~optionally substituted pyrrolidinyl, optionally substituted piperazinyl, optionally substituted piperidinyl, morpholinyl, 1,2,3,6-tetrahydro-pyridinyl, (optionally substituted pyrrolidinyl) C₁-C₂-alkyl, (optionally substituted piperidinyl) C₁-C₂-alkyl, (optionally substituted piperazinyl) C₁-C₂-alkyl, morpholinyl C₁-C₂-alkyl, C₁-C₄-alkylamino-C₁-C₄-alkyl, C₁-C₄-hydroxyalkylamino, (optionally substituted pyrrolidinyl) C₁-C₂-alkylamino, (optionally substituted piperidinyl) C₁-C₂-alkylamino, (optionally substituted piperazinyl) C₁-C₂-alkylamino, morpholinyl C₁-C₂-alkylamino, optionally substituted pyrrolidinyl-C₁-C₄-alkoxy, optionally substituted azetidiny-C₁-C₄-alkoxy, tetrahydrofuryl-C₁-C₄-alkoxy, optionally substituted piperidinyl-C₁-C₄-alkoxy, C₁-C₄-alkylamino-C₁-C₄-alkoxy, tetrahydrofuryloxy, optionally substituted piperidinyl, and optionally substituted phenoxy, C₁-C₄-alkylaminocarbonyl and C₁-C₄-alkylaminothiocarbonyl;~~
wherein R¹⁶ is selected from H, ~~5-6-membered nitrogen-containing heterocyclylcarboxyl, C₁-C₄-alkylaminocarbonyl, and C₁-C₄-alkylaminomethyl, and 5-6-membered nitrogen-containing heterocyclylmethyl;~~ and wherein R¹⁷ is selected from halo, C₁-C₂-alkyl, ~~optionally substituted 5-6-membered heterocyclylsulfonyl C₁-C₂-alkyl, optionally substituted phenoxy, and C₃-C₆-cycloalkyl-C₂-C₄-alkynyl, and pharmaceutically acceptable derivatives thereof.~~

Claim 125 (currently amended): The method of Claim 124 wherein R¹⁵ is selected from ~~H,~~
tetrahydro-furanyloxy, 1-methylpyrrolidin-2-ylmethoxy, 2-pyrrolidinylmethoxy, 3-

pyrrolidinylmethoxy, 1-Boc-pyrrolidin-2-ylmethoxy, 4-piperidinylmethoxy, 1-Boc-piperidin-4-ylmethoxy, 1-Boc-piperidin-4-ylethoxy, piperidin-4-ylethoxy, 1-methyl-piperidin-4-ylmethoxy, 1-Boc-azetidin-3-ylmethoxy, 1-methyl-azetidin-3-ylmethoxy, 3-azetidylmethoxy, 1-methyl-piperidin-4-yloxy, ~~phenyloxy, phenoxy, 4-(pyrrolidin-1-ylmethyl)phenoxy, and~~
~~dimethylaminoethoxy, 1-piperidinylmethyl, 1-(piperidin-1-yl)ethyl, 3-methylpiperidin-1-ylmethyl, 1-pyrrolidinylmethyl, 2,2,6,6-tetramethylpiperidin-1-ylmethyl, 2,6-dimethylpiperidin-1-ylmethyl, dimethylaminomethyl, diethylaminomethyl, diethylaminothiocarbonyl, diethylaminocarbonyl, N-Boc-N-isopropylaminomethyl, isopropylaminomethyl, 2-thienylsulfonylmethyl, hydroxypropylamine, 4-ethyl-piperidin-1-yl, 4-(2-pyridyl)piperidin-1-yl, 1-methylpiperidin-4-yl, 4-(2-pyrazinyl)piperidin-1-yl, 1-methyl-1,2,3,6-tetrahydro-pyridin-4-yl, 1,2,3,6-tetrahydro-pyridin-4-yl, and 1-Boc-1,2,3,6-tetrahydro-pyridin-4-yl; wherein R¹⁶ is selected from H, 1-piperidinylcarbonyl, diethylaminocarbonyl, and diethylaminomethyl, 1-piperidinylmethyl; and wherein R¹⁷ is selected from chloro, bromo, methyl and cyclopropylethynyl; and pharmaceutically acceptable derivatives thereof.~~

Claim 126 (currently amended): The method of Claim 125, wherein R¹⁷ is chloro or bromo; ~~and pharmaceutically acceptable derivatives thereof.~~

Claim 127 (currently amended): The method of Claim 123, wherein R¹⁵ is selected from H, ~~optionally substituted pyrrolidinyl, optionally substituted piperazinyl, optionally substituted piperidinyl, morpholinyl, 1,2,3,6-tetrahydro-pyridinyl, (optionally substituted pyrrolidinyl)-C₁-C₂-alkyl, (optionally substituted piperidinyl)-C₁-C₂-alkyl, (optionally substituted piperazinyl)-C₁-C₂-alkyl, morpholinyl-C₁-C₂-alkyl, (optionally substituted pyrrolidinyl)-C₁-C₂-alkylamine, (optionally substituted piperidinyl)-C₁-C₂-alkylamine, (optionally substituted piperazinyl)-C₁-C₂-alkylamine, morpholinyl-C₁-C₂-alkylamine, C₁-C₄-alkylamine-C₁-C₄-alkyl, C₁-C₄-hydroxyalkylamine, optionally substituted pyrrolidinyl-C₁-C₄-alkoxy, optionally substituted azetidyl-C₁-C₄-alkoxy, tetrahydrofuryl-C₁-C₄-alkoxy, optionally substituted piperidinyl-C₁-C₄-alkoxy, C₁-C₄-alkylamino-C₁-C₄-alkoxy, tetrahydrofuryloxy, optionally substituted piperidinyl, and optionally substituted phenoxy, C₁-C₄-alkylaminocarbonyl and C₁-C₄-alkylaminothiocarbonyl; wherein R¹⁶ is selected from H, 5-6-membered nitrogen-containing heterocyclylcarbonyl, C₁-C₄-alkylaminocarbonyl, and C₁-C₄-alkylaminomethyl; and 5-6-~~

~~membered nitrogen-containing heterocyclylmethyl; and wherein R¹⁷ is selected from C₃-C₆-cycloalkyl and phenyl- optionally substituted with one or two substituents selected from halo, C₁-C₄-alkylamino, amino, nitro, C₁-C₄-alkoxy, C₁-C₂-haloalkyl, hydroxy, C₁-C₄-alkylthio, C₁-C₄-alkylcarbonylamino, (optionally substituted phenyl)sulfonylamino, cyano, C₁-C₂-haloalkoxy, 5- or 6- membered N-containing heterocyclyl, aminosulfonyl, (6- membered N-containing heterocyclyl)sulfonyl, C₁-C₂-haloalkylcarbonylaminesulfonyl and (optionally substituted phenyl)aminosulfonyl; and pharmaceutically acceptable derivatives thereof.~~

Claim 128 (currently amended): The method of Claim 127, wherein R¹⁵ is selected from H, tetrahydro-furanyloxy, 1-methylpyrrolidin-2-ylmethoxy, 2-pyrrolidinylmethoxy, 3-pyrrolidinylmethoxy, 1-Boc-pyrrolidin-2-ylmethoxy, 4-piperidinylmethoxy, 1-Boc-piperidin-4-ylmethoxy, 1-Boc-piperidin-4-ylethoxy, piperidin-4-ylethoxy, 1-methyl-piperidin-4-ylmethoxy, 1-Boc-azetidin-3-ylmethoxy, 1-methyl-azetidin-3-ylmethoxy, 3-azetidinylmethoxy, 1-methyl-piperidin-4-yloxy, ~~phenyloxy~~, phenoxy 4-(pyrrolidin-1-ylmethyl)phenoxy, and dimethylaminoethoxy, ~~1-piperidinylmethyl, 1-(piperidin-1-yl)ethyl, 3-methylpiperidin-1-ylmethyl, 1-pyrrolidinylmethyl, 2,2,6,6-tetramethylpiperidin-1-ylmethyl, 2,6-dimethylpiperidin-1-ylmethyl, dimethylaminomethyl, diethylaminomethyl, diethylaminothiocarbonyl, diethylaminocarbonyl, N-Boc-N-isopropylaminomethyl, isopropylaminomethyl, 2-thienylsulfonylmethyl, hydroxypropylamino, 4-ethyl-piperidin-1-yl, 4-(2-pyridyl)piperidin-1-yl, 1-methylpiperidin-4-yl, 4-(2-pyrazinyl)piperidin-1-yl, 1-methyl-1,2,3,6-tetrahydro-pyridin-4-yl, 1,2,3,6-tetrahydro-pyridin-4-yl, and 1-Boc-1,2,3,6-tetrahydro-pyridin-4-yl~~; wherein R¹⁶ is selected from H, ~~1-piperidinylcarbonyl~~, diethylaminocarbonyl, and diethylaminomethyl, ~~1-piperidinylmethyl~~; and wherein R¹⁷ is selected from cyclopropyl and phenyl-~~optionally substituted with aminosulfonyl; and pharmaceutically acceptable derivatives thereof.~~

Claim 129 (currently amended): The method of Claim 128, wherein R¹⁷ is unsubstituted phenyl; ~~and pharmaceutically acceptable derivatives thereof.~~

Claim 130 (currently amended): The method of Claim 123, wherein R¹⁵ is selected from H, ~~optionally substituted pyrrolidinyl, optionally substituted piperazinyl, optionally substituted piperidinyl, morpholinyl, 1,2,3,6-tetrahydro-pyridinyl, (optionally substituted pyrrolidinyl) C₁-~~

~~C₂-alkyl, (optionally substituted piperidinyl) C₄-C₂-alkyl, (optionally substituted piperazinyl) C₄-C₂-alkyl, morpholinyl C₄-C₂-alkyl, (optionally substituted pyrrolidinyl) C₄-C₂-alkylamino, (optionally substituted piperidinyl) C₄-C₂-alkylamino, (optionally substituted piperazinyl) C₄-C₂-alkylamino, morpholinyl C₄-C₂-alkylamino, C₄-C₄-alkylamino C₄-C₄-alkyl, C₄-C₄-hydroxyalkylamino, optionally substituted pyrrolidinyl-C₁-C₄-alkoxy, optionally substituted azetidiny-C₁-C₄-alkoxy, tetrahydrofuryl-C₁-C₄-alkoxy, optionally substituted piperidinyl-C₁-C₄-alkoxy, C₁-C₄-alkylamino-C₁-C₄-alkoxy, tetrahydrofuryloxy, optionally substituted piperidinyl, and optionally substituted phenoxy, C₄-C₄-alkylaminocarbonyl and C₄-C₄-alkylaminethiocarbonyl; wherein R¹⁶ is selected from H, 5-6 membered nitrogen-containing heterocyclylcarbonyl, C₁-C₄-alkylaminocarbonyl, and C₁-C₄-alkylaminomethyl, and 5-6 membered nitrogen-containing heterocyclylmethyl; and wherein R¹⁷ is selected from optionally substituted indazolyl, optionally substituted indolyl, unsubstituted 5-membered oxygen or sulfur containing heterocaryl, unsubstituted thienyl, unsubstituted 6-membered nitrogen-containing heterocyclyl, and 6-membered nitrogen-containing heterocyclyl substituted with one or more substituents independently selected from pyridyl, phenyl,~~

~~C₄-C₄-alkyl, C₄-C₂-haloalkyl, C₁-C₂ alkoxy, amino, halo, piperidinyl, morpholinyl, C₄-C₂-alkylpiperazinyl, C₄-C₂-alkylaminethiocarbonyl, N,N-di C₄-C₂-alkylamino C₄-C₄-alkylenyl, N-C₄-C₂-alkylamino C₄-C₄-alkylenyl, morpholinyl C₄-C₄-alkylenylaminocarbonyl, aminocarbonyl, C₄-C₂-haloalkylcarbonylamino, morpholinyl C₄-C₄-alkylenylamino, N,N-di C₄-C₂-alkylamino and N,N-di C₄-C₂-alkylamino C₄-C₄-alkylenylamino;~~

~~and pharmaceutically acceptable derivatives thereof.~~

Claim 131 (currently amended): The method of Claim 130, wherein R¹⁵ is selected from H, tetrahydro-furanyloxy, 1-methylpyrrolidin-2-ylmethoxy, 2-pyrrolidinylmethoxy, 3-pyrrolidinylmethoxy, 1-Boc-pyrrolidin-2-ylmethoxy, 4-piperidinylmethoxy, 1-Boc-piperidin-4-ylmethoxy, 1-Boc-piperidin-4-ylethoxy, piperidin-4-ylethoxy, 1-methyl-piperidin-4-ylmethoxy, 1-Boc-azetidin-3-ylmethoxy, 1-methyl-azetidin-3-ylmethoxy, 3-azetidylmethoxy, 1-methyl-piperidin-4-yloxy, phenyloxy, 4-(pyrrolidin-1-ylmethyl)phenoxy, and dimethylaminoethoxy, 1-piperidinylmethyl, 1-(piperidin-1-yl)ethyl, 3-methylpiperidin-1-ylmethyl, 1-pyrrolidinylmethyl, 2,2,6,6-tetramethylpiperidin-1-ylmethyl, 2,6-dimethylpiperidin-1-ylmethyl, dimethylaminomethyl, diethylaminomethyl, diethylaminethiocarbonyl, diethylaminocarbonyl, N-

~~Boc-N-isopropylaminomethyl, isopropylaminomethyl, 2-thienylsulfenylmethyl, hydroxypropylamino, 4-ethyl-piperidin-1-yl, 4-(2-pyridyl)piperidin-1-yl, 1-methylpiperidin-4-yl, 4-(2-pyrazinyl)piperidin-1-yl, 1-methyl-1,2,3,6-tetrahydro-pyridin-4-yl, 1,2,3,6-tetrahydro-pyridin-4-yl, and 1-Boc-1,2,3,6-tetrahydro-pyridin-4-yl; wherein R¹⁶ is selected from H, 1-piperidinylcarbonyl, diethylaminocarbonyl, and diethylaminomethyl, 1-piperidinylmethyl; and wherein R¹⁷ is selected from 5-indazolyl, 1-Boc-indol-5-yl, unsubstituted thienyl, 5-tert-butylloxazol-2-yl and 4-pyridyl substituted with one or more substituents independently selected from methoxy and chloro; and pharmaceutically acceptable derivatives thereof.~~

Claim 132 (currently amended): The method of Claim ~~130~~ 123, wherein R¹⁷ is 4-pyridyl, ~~and pharmaceutically acceptable derivatives thereof.~~

Claim 133 (currently amended): The method of Claim 123, wherein the compound is ~~and pharmaceutically acceptable derivatives thereof~~ selected from:

~~1-[6-(3-Methyl-piperidin-1-ylmethyl)-pyridin-2-yl]-3-(2-phenyl-thiazol-4-yl)-urea;
1-[4-(Piperidine-1-carbonyl)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;
1-(2-Chloro-thiazol-4-yl)-3-[4-(piperidine-1-carbonyl)-pyridin-2-yl]-urea;
N,N-Diethyl-2-[3-(2-pyridin-4-yl-thiazol-4-yl)-ureido]-isonicotinamide;
N,N-Diethyl-2-[3-(2-phenyl-thiazol-4-yl)-ureido]-isonicotinamide;
2-[3-(2-Bromo-thiazol-4-yl)-ureido]-N,N-diethyl-isonicotinamide;
1-(4-Diethylaminomethyl-pyridin-2-yl)-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;
1-[6-(2,6-Dimethyl-piperidin-1-ylmethyl)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;
1-[6-(1-Piperidin-1-yl-ethyl)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;
2-((6-[3-(2-Pyridin-4-yl-thiazol-4-yl)-ureido]-pyridin-2-ylamino)-methyl)-piperidine-1-carboxylic acid tert-butyl ester;
1-[6-[(Piperidin-2-ylmethyl)-amino]-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;
(S)-1-[6-(3-Methyl-piperidin-1-ylmethyl)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;
(R)-1-[6-(3-Methyl-piperidin-1-ylmethyl)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;
1-(2-Chloro-thiazol-4-yl)-3-(6-piperidin-1-ylmethyl-pyridin-2-yl)-urea;
1-(2-Bromo-thiazol-4-yl)-3-[6-(2-piperidin-4-yl-ethoxy)-pyridin-2-yl]-urea;~~

1-(2-Chloro-thiazol-4-yl)-3-[6-(2-piperidin-4-yl-ethoxy)-pyridin-2-yl]-urea;
1-[6-(Azetidin-3-ylmethoxy)-pyridin-2-yl]-3-(2-bromo-thiazol-4-yl)-urea;
1-[6-(Azetidin-3-ylmethoxy)-pyridin-2-yl]-3-(2-chloro-thiazol-4-yl)-urea;
1-(2-Bromo-thiazol-4-yl)-3-[6-(piperidin-4-ylmethoxy)-pyridin-2-yl]-urea;
1-(2-Chloro-thiazol-4-yl)-3-[6-(piperidin-4-ylmethoxy)-pyridin-2-yl]-urea;
tert-Butyl 3-{6-[3-(2-pyridin-4-yl-thiazol-4-yl)-ureido]-pyridin-2-yloxymethyl}-pyrrolidine-1-carboxylate;
1-(2-Pyridin-4-yl-thiazol-4-yl)-3-[6-(pyrrolidin-3-ylmethoxy)-pyridin-2-yl]-urea;
1-(2-Cyclopropyl-thiazol-4-yl)-3-[6-(2-piperidin-4-yl-ethoxy)-pyridin-2-yl]-urea;
~~1-[6-(Isopropylamine methyl) pyridin 2-yl] 3-(2-pyridin-4-yl-thiazol-4-yl)-urea;~~
~~1-[6-(Isopropylamine methyl) pyridin 2-yl] 3-(2-phenyl-thiazol-4-yl)-urea;~~
~~1-(2-Bromo-thiazol-4-yl)-3-[6-(isopropylamine methyl)-pyridin-2-yl]-urea;~~
1-(2-Bromo-thiazol-4-yl)-3-[6-(1-methyl-pyrrolidin-2-ylmethoxy)-pyridin-2-yl]-urea;
1-(2-Chloro-thiazol-4-yl)-3-[6-(1-methyl-pyrrolidin-2-ylmethoxy)-pyridin-2-yl]-urea;
1-(2-phenylthiazol-4-yl)-3-(6-*p*-pyrrolidin-1-ylmethylphenoxy)pyridin-2-yl)urea;
1-(2-Pyridin-4-yl-thiazol-4-yl)-3-[6-(tetrahydro-furan-3-yloxy)-pyridin-2-yl]-urea;
~~1-[2-(1H-Indazol-5-yl) thiazol-4-yl] 3-(6-piperidin-1-ylmethyl pyridin-2-yl)-urea;~~
~~1-(1'-Methyl-1',2',3',6'-tetrahydro-[2,4']bipyridinyl-6-yl) 3-(2-pyridin-4-yl-thiazol-4-yl)-urea;~~
~~1-(2-Bromo-thiazol-4-yl) 3-(1'-methyl-1',2',3',6'-tetrahydro-[2,4']bipyridinyl-6-yl)-urea;~~
~~1-(1'-Methyl-1',2',3',6'-tetrahydro-2[2,4]bipyridinyl-6-yl) 3-(2-phenyl-thiazol-4-yl)-urea;~~
~~1-[6-(3-Hydroxy-propylamine) pyridin 2-yl] 3-(2-pyridin-4-yl-thiazol-4-yl)-urea;~~
~~1-(2-Bromo-thiazol-4-yl) 3-[6-(3-hydroxy-propylamine) pyridin-2-yl]-urea;~~
~~1-(1'-Methyl-1',2',3',4',5',6'-hexahydro-[2,4']bipyridinyl-6-yl) 3-(2-pyridin-4-yl-thiazol-4-yl)-urea;~~
~~urea;~~
~~1-(1'-Methyl-1',2',3',4',5',6'-hexahydro-[2,4']bipyridinyl-6-yl) 3-(2-phenyl-thiazol-4-yl)-urea;~~
~~6-[3-(2-Pyridin-4-yl-thiazol-4-yl)-ureido]-3',6'-dihydro-2'H-[2,4]bipyridinyl-1'-carboxylic acid~~
~~*tert*-butylester;~~
~~1-(2-Pyridin-4-yl-thiazol-4-yl) 3-(1',2',3',6'-tetrahydro-[2,4']bipyridinyl-6-yl)-urea;~~
1-(2-Pyridin-4-yl-thiazol-4-yl)-3-[6-(tetrahydro-furan-3-ylmethoxy)-pyridin-2-yl]-urea;

2-[6-[3-(2-Pyridin-4-yl-thiazol-4-yl)-ureido]-pyridin-2-yloxymethyl]-pyrrolidine-1-carboxylic acid tert-butyl ester;

1-(2-Pyridin-4-yl-thiazol-4-yl)-3-[6-(pyrrolidin-2-ylmethoxy)-pyridin-2-yl]-urea;

~~6-[3-(2-Pyridin-4-yl-thiazol-4-yl)-ureido]-pyridine-2-carbothioic acid diethylamide;~~

~~1-(2-Bromo-thiazol-4-yl)-3-[6-(3-methyl-piperidin-1-ylmethyl)-pyridin-2-yl]-urea;~~

~~1-(2-Chloro-thiazol-4-yl)-3-[6-(3-methyl-piperidin-1-ylmethyl)-pyridin-2-yl]-urea;~~

~~1-(2-Phenyl-thiazol-4-yl)-3-[4-(piperidine-1-carbonyl)-pyridin-2-yl]-urea;~~

~~1-(2-Bromo-thiazol-4-yl)-3-[4-(piperidine-1-carbonyl)-pyridin-2-yl]-urea;~~

1-[2-(2-Methoxy-pyridin-4-yl)-thiazol-4-yl]-3-(6-phenoxy-pyridin-2-yl)-urea;

1-[2-(2-Methoxy-pyridin-4-yl)-thiazol-4-yl]-3-[6-(1-methyl-pyrrolidin-2-ylmethoxy)-pyridin-2-yl]-urea;

1-[6-(2-Dimethylamino-ethoxy)-pyridin-2-yl]-3-[2-(2-methoxy-pyridin-4-yl)-thiazol-4-yl]-urea;

1-[6-(1-Methyl-pyrrolidin-2-ylmethoxy)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;

~~1-(2-phenylthiazol-4-yl)-3-(6-pyrrolidin-1-ylmethyl-pyridin-2-yl)urea;~~

~~1-(6-Diethylaminomethylpyridin-2-yl)-3-(2-phenylthiazol-4-yl)urea;~~

(S)-1-[6-(1-Methylpyrrolidin-2-ylmethoxy)pyridin-2-yl]-3-(2-phenylthiazol-4-yl)urea;

1-[6-(2-Piperidin-4-yl-ethoxy)pyridin-2-yl]-3-[2-phenylthiazol-4-yl]urea;

~~1-[6-(4-Ethylpiperazin-1-yl)-pyridin-2-yl]-3-(2-phenylthiazol-4-yl)urea;~~

~~Diethyl 6-[3-(2-phenylthiazol-4-yl)ureido]-pyridine-2-carboxamide;~~

1-(2-Pyridin-4-yl-thiazol-4-yl)-3-(6-*p*-pyrrolidin-1-ylmethylphenoxy)pyridin-2-yl)urea;

1-(2-Bromothiazol-4-yl)-3-(6-*p*-pyrrolidin-1-ylmethylphenoxy)pyridin-2-yl)urea;

1-[6-(Piperidin-4-ylmethoxy)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;

1-[6-(1-Methyl-piperidin-4-ylmethoxy)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;

1-[6-(1-Methyl-piperidin-4-yloxy)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;

1-[6-(1-Methyl-azetidin-3-ylmethoxy)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;

1-[6-(Azetidin-3-ylmethoxy)-pyridin-2-yl]-3-(2-phenyl-thiazol-4-yl)-urea;

1-[6-(1-Methyl-azetidin-3-ylmethoxy)-pyridin-2-yl]-3-(2-phenyl-thiazol-4-yl)-urea;

1-(2-Phenyl-thiazol-4-yl)-3-[6-(piperidin-4-ylmethoxy)-pyridin-2-yl]-urea;

1-[6-(1-Methyl-piperidin-4-ylmethoxy)-pyridin-2-yl]-3-(2-phenyl-thiazol-4-yl)-urea;

1-[6-(1-Methyl-piperidin-4-yloxy)-pyridin-2-yl]-3-(2-phenyl-thiazol-4-yl)-urea;

1-[6-(2-Piperidin-4-yl-ethoxy)-pyridin-2-yl]-3-(2-thiophen-2-yl-thiazol-4-yl)-urea; and

1-[6-(1-Methyl-pyrrolidin-2-ylmethoxy)-pyridin-2-yl]-3-[2-(thiophene-2-sulfonylmethyl)-thiazol-4-yl]-urea;

~~1-[2-(2-Methoxy-pyridin-4-yl)-thiazol-4-yl]-3-(6-piperidin-1-ylmethyl-pyridin-2-yl)-urea; and~~
~~[2-(2-Chloro-pyridin-4-yl)-thiazol-4-yl]-3-(6-piperidin-1-ylmethyl-pyridin-2-yl)-urea;~~
and pharmaceutically acceptable salts thereof.

Claim 134 (currently amended): The method of Claim 123, wherein the compound is ~~and~~
~~pharmaceutically acceptable derivatives thereof~~ selected from:

~~1-[6-(3-Methyl-piperidin-1-ylmethyl)-pyridin-2-yl]-3-(2-phenyl-thiazol-4-yl)-urea;~~
~~1-[4-(Piperidine-1-carbonyl)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;~~
~~N,N-Diethyl-2-[3-(2-pyridin-4-yl-thiazol-4-yl)-ureido]-isonicotinamide;~~
~~1-(4-Diethylaminomethyl-pyridin-2-yl)-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;~~
~~1-[6-(2,6-Dimethyl-piperidin-1-ylmethyl)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;~~
~~1-[6-(1-Piperidin-1-yl-ethyl)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;~~
~~2-((6-[3-(2-Pyridin-4-yl-thiazol-4-yl)-ureido]-pyridin-2-ylamino)-methyl)-piperidine-1-carboxylic acid tert-butyl ester;~~
~~1-{6-[(Piperidin-2-ylmethyl)-amino]-pyridin-2-yl}-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;~~
~~(S)-1-[6-(3-Methyl-piperidin-1-ylmethyl)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;~~
~~(R)-1-[6-(3-Methyl-piperidin-1-ylmethyl)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;~~
~~1-(2-Chloro-thiazol-4-yl)-3-(6-piperidin-1-ylmethyl-pyridin-2-yl)-urea;~~
1-(2-Bromo-thiazol-4-yl)-3-[6-(2-piperidin-4-yl-ethoxy)-pyridin-2-yl]-urea;
1-(2-Chloro-thiazol-4-yl)-3-[6-(2-piperidin-4-yl-ethoxy)-pyridin-2-yl]-urea;
1-[6-(Azetidin-3-ylmethoxy)-pyridin-2-yl]-3-(2-bromo-thiazol-4-yl)-urea;
1-[6-(Azetidin-3-ylmethoxy)-pyridin-2-yl]-3-(2-chloro-thiazol-4-yl)-urea;
1-(2-Bromo-thiazol-4-yl)-3-[6-(piperidin-4-ylmethoxy)-pyridin-2-yl]-urea;
1-(2-Chloro-thiazol-4-yl)-3-[6-(piperidin-4-ylmethoxy)-pyridin-2-yl]-urea;
3-(4-{3-[6-(1-Methyl-pyrrolidin-2-ylmethoxy)-pyridin-2-yl]-ureido}-thiazol-2-yl)-
benzenesulfonamide;
tert-Butyl 3-{6-[3-(2-pyridin-4-yl-thiazol-4-yl)-ureido]-pyridin-2-yloxymethyl}-pyrrolidine-1-
carboxylate;
1-(2-Pyridin-4-yl-thiazol-4-yl)-3-[6-(pyrrolidin-3-ylmethoxy)-pyridin-2-yl]-urea;

1-(2-Cyclopropyl-thiazol-4-yl)-3-[6-(2-piperidin-4-yl-ethoxy)-pyridin-2-yl]-urea;
~~Isopropyl {6 [3 (2 pyridin 4 yl thiazol 4 yl) ureido] pyridin 2 ylmethyl} carbamic acid tert-butyl ester;~~
~~1 [6 (Isopropylamino methyl) pyridin 2 yl] 3 (2 pyridin 4 yl thiazol 4 yl) urea;~~
~~Isopropyl {6 [3 (2 phenyl thiazol 4 yl) ureido] pyridin 2 ylmethyl} carbamic acid tert-butyl ester;~~
~~1 [6 (Isopropylamino methyl) pyridin 2 yl] 3 (2 phenyl thiazol 4 yl) urea;~~
1-(2-Bromo-thiazol-4-yl)-3-[6-(1-methyl-pyrrolidin-2-ylmethoxy)-pyridin-2-yl]-urea;
1-(2-Chloro-thiazol-4-yl)-3-[6-(1-methyl-pyrrolidin-2-ylmethoxy)-pyridin-2-yl]-urea;
1-(2-phenylthiazol-4-yl)-3-(6-*p*-pyrrolidin-1-ylmethylphenoxy)pyridin-2-yl)urea;
1-(2-Pyridin-4-yl-thiazol-4-yl)-3-[6-(tetrahydro-furan-3-yloxy)-pyridin-2-yl]-urea;
~~1 [2 (1H Indazol 5 yl) thiazol 4 yl] 3 (6 piperidin 1 ylmethyl pyridin 2 yl) urea;~~
~~1 (1' Methyl 1',2',3',6' tetrahydro [2,4']bipyridinyl 6 yl) 3 (2 pyridin 4 yl thiazol 4 yl) urea;~~
~~1 (2 Bromo thiazol 4 yl) 3 (1' methyl 1',2',3',6' tetrahydro [2,4']bipyridinyl 6 yl) urea;~~
~~1 (1' Methyl 1',2',3',6' tetrahydro 2[2,4]bipyridinyl 6 yl) 3 (2 phenyl thiazol 4 yl) urea;~~
~~1 [6 (3 Hydroxy propylamino) pyridin 2 yl] 3 (2 pyridin 4 yl thiazol 4 yl) urea;~~
~~1 (2 Bromo thiazol 4 yl) 3 [6(3 hydroxy propylamino) pyridin 2 yl] urea;~~
~~1 (1' Methyl 1',2',3',4',5',6' hexahydro [2,4']bipyridinyl 6 yl) 3 (2 pyridin 4 yl thiazol 4 yl) urea;~~
~~1 (1' Methyl 1',2',3',4',5',6' hexahydro [2,4']bipyridinyl 6 yl) 3 (2 phenyl thiazol 4 yl) urea;~~
~~6 [3 (2 Pyridin 4 yl thiazol 4 yl) ureido] 3',6' dihydro 2'H [2,4]bipyridinyl 1' carboxylic acid tert-butylester;~~
~~1 (2 Pyridin 4 yl thiazol 4 yl) 3 (1',2',3',6' tetrahydro [2,4']bipyridinyl 6 yl) urea;~~
1-(2-Pyridin-4-yl-thiazol-4-yl)-3-[6-(tetrahydro-furan-3-ylmethoxy)-pyridin-2-yl]-urea;
2-[6-[3-(2-Pyridin-4-yl-thiazol-4-yl)-ureido]-pyridin-2-yloxymethyl]-pyrrolidine-1-carboxylic acid tert-butyl ester;
1-(2-Pyridin-4-yl-thiazol-4-yl)-3-[6-(pyrrolidin-2-ylmethoxy)-pyridin-2-yl]-urea;
~~6 [3 (2 Pyridin 4 yl thiazol 4 yl) ureido] pyridine 2 carbothioic acid diethylamide;~~
~~1 (2 Bromo thiazol 4 yl) 3 [6 (3 methyl piperidin 1 ylmethyl) pyridin 2 yl] urea;~~
~~1 (2 Chloro thiazol 4 yl) 3 [6 (3 methyl piperidin 1 ylmethyl) pyridin 2 yl] urea;~~
1-[2-(2-Methoxy-pyridin-4-yl)-thiazol-4-yl]-3-[6-(1-methyl-pyrrolidin-2-ylmethoxy)-pyridin-2-yl]-urea;

1-[6-(2-Dimethylamino-ethoxy)-pyridin-2-yl]-3-[2-(2-methoxy-pyridin-4-yl)-thiazol-4-yl]-urea;
1-[6-(1-Methyl-pyrrolidin-2-ylmethoxy)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;
~~1-(2-phenylthiazol-4-yl)-3-(6-pyrrolidin-1-ylmethyl-pyridin-2-yl)urea;~~
~~1-(6-Diethylaminomethylpyridin-2-yl)-3-(2-phenylthiazol-4-yl)urea;~~
(S)-1-[6-(1-Methylpyrrolidin-2-ylmethoxy)pyridin-2-yl]-3-(2-phenylthiazol-4-yl)urea;
1-[6-(2-Piperidin-4-yl-ethoxy)pyridin-2-yl]-3-[2-phenylthiazol-4-yl]urea;
~~1-[6-(4-Ethylpiperazin-1-yl)-pyridin-2-yl]-3-(2-phenylthiazol-4-yl)urea;~~
~~1-(2-phenylthiazol-4-yl)-3-[6-(4-pyrimidin-2-yl-piperazin-1-yl)pyridin-2-yl]urea;~~
~~Diethyl 6-[3-(2-phenylthiazol-4-yl)ureido]pyridine-2-carboxamide;~~
1-(2-Pyridin-4-yl-thiazol-4-yl)-3-(6-*p*-pyrrolidin-1-ylmethylphenoxy)pyridin-2-yl)urea;
1-(2-Bromothiazol-4-yl)-3-(6-*p*-pyrrolidin-1-ylmethylphenoxy)pyridin-2-yl)urea;
1-[6-(Piperidin-4-ylmethoxy)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;
1-[6-(1-Methyl-piperidin-4-ylmethoxy)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;
1-[6-(1-Methyl-piperidin-4-yloxy)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;
1-[6-(1-Methyl-azetidin-3-ylmethoxy)-pyridin-2-yl]-3-(2-pyridin-4-yl-thiazol-4-yl)-urea;
1-[6-(Azetidin-3-ylmethoxy)-pyridin-2-yl]-3-(2-phenyl-thiazol-4-yl)-urea;
1-[6-(1-Methyl-azetidin-3-ylmethoxy)-pyridin-2-yl]-3-(2-phenyl-thiazol-4-yl)-urea;
1-(2-Phenyl-thiazol-4-yl)-3-[6-(piperidin-4-ylmethoxy)-pyridin-2-yl]-urea;
1-[6-(1-Methyl-piperidin-4-ylmethoxy)-pyridin-2-yl]-3-(2-phenyl-thiazol-4-yl)-urea;
1-[6-(1-Methyl-piperidin-4-yloxy)-pyridin-2-yl]-3-(2-phenyl-thiazol-4-yl)-urea;
1-[6-(2-Piperidin-4-yl-ethoxy)-pyridin-2-yl]-3-(2-thiophen-2-yl-thiazol-4-yl)-urea; and
1-[6-(1-Methyl-pyrrolidin-2-ylmethoxy)-pyridin-2-yl]-3-[2-(thiophene-2-sulfonylmethyl)-thiazol-4-yl]-urea;
~~1-[2-(2-Methoxy-pyridin-4-yl)-thiazol-4-yl]-3-(6-piperidin-1-ylmethyl-pyridin-2-yl)-urea; and~~
~~[2-(2-Chloro-pyridin-4-yl)-thiazol-4-yl]-3-(6-piperidin-1-ylmethyl-pyridin-2-yl)-urea;~~
and pharmaceutically acceptable salts thereof.